

10/141,986

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STRUCTURE FILE UPDATES: 2 JUL 2008 HIGHEST RN 1032450-11-5
DICTIONARY FILE UPDATES: 2 JUL 2008 HIGHEST RN 1032450-11-5

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TSCA INFORMATION NOW CURRENT THROUGH January 9, 2008.

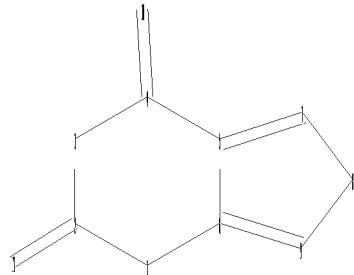
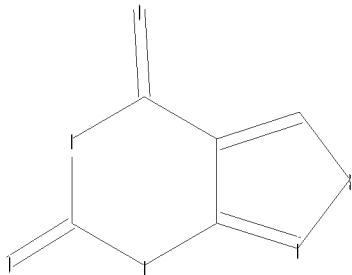
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<http://www.cas.org/support/stngen/stndoc/properties.html>

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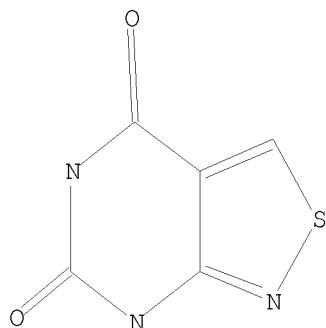


chain nodes :
10 11
ring nodes :
1 2 3 4 5 6 7 8 9
chain bonds :
2-11 4-10
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9
exact/norm bonds :
1-2 1-6 2-3 2-11 3-4 4-5 4-10 5-6 5-7 6-9 7-8 8-9

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS

L1 STRUCTURE UPLOADED

=> d
L1 HAS NO ANSWERS
L1 STR



Structure attributes must be viewed using STN Express query preparation.

10/141,986

=> s 11 sam
SAMPLE SEARCH INITIATED 15:44:46 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 30 TO ITERATE

100.0% PROCESSED 30 ITERATIONS 3 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 272 TO 928
PROJECTED ANSWERS: 3 TO 163

L2 3 SEA SSS SAM L1

=> s 11 ful
FULL SEARCH INITIATED 15:44:53 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 606 TO ITERATE

100.0% PROCESSED 606 ITERATIONS 111 ANSWERS
SEARCH TIME: 00.00.01

L3 111 SEA SSS FUL L1

=> fil cap1
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FILE COVERS 1907 - 3 Jul 2008 VOL 149 ISS 1
FILE LAST UPDATED: 2 Jul 2008 (20080702/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

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=> s 13

10/141,986

L4 24 L3

=> s 14 not (2008so or 2007/so or 2006/so or 2005/so)

0 2008SO

941580 2007/SO

943642 2006/SO

884541 2005/SO

L5 23 L4 NOT (2008SO OR 2007/SO OR 2006/SO OR 2005/SO)

=> s 14 not (2008/so or 2007/so or 2006/so or 2005/so)

402651 2008/SO

941580 2007/SO

943642 2006/SO

884541 2005/SO

L6 23 L4 NOT (2008/SO OR 2007/SO OR 2006/SO OR 2005/SO)

=> d 16 ibib hitstr abs 1-23

L6 ANSWER 1 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:902901 CAPLUS
 DOCUMENT NUMBER: 143:229880
 TITLE: Preparation of isothiazolopyrimidinediones as parathyroid hormone type I receptor (PTH1R) agonists for treatment of osteoporosis.
 INVENTOR(S): Spearing, Paul Kenneth
 PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
 SOURCE: PCT Int. Appl., 21 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005077959	A1	20050825	WO 2005-US3247	20050203
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1713815	A1	20061025	EP 2005-712622	20050203
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, HR, IS				
JP 2007522214	T	20070809	JP 2006-553161	20050203
US 20070099940	A1	20070503	US 2006-597860	20060810
PRIORITY APPLN. INFO.:			US 2004-543764P	P 20040211
			WO 2005-US3247	W 20050203

OTHER SOURCE(S): CASREACT 143:229880; MARPAT 143:229880

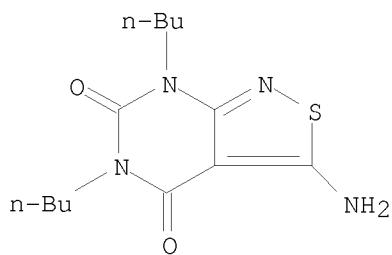
IT 52903-32-9P 862980-97-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(claimed compound; preparation of isothiazolopyrimidinediones as parathyroid hormone type I receptor agonists for treatment of osteoporosis)

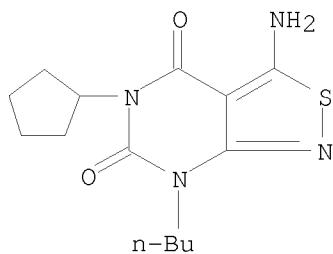
RN 52903-32-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dibutyl- (CA INDEX NAME)



RN 862980-97-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-butyl-5-cyclopentyl- (CA INDEX NAME)



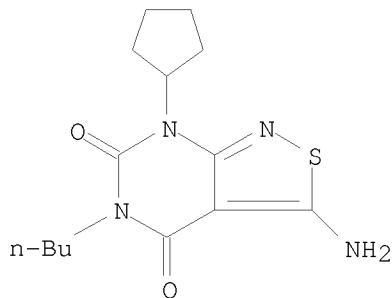
IT 862981-03-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

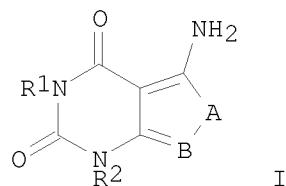
(preparation of isothiazolopyrimidinediones as parathyroid hormone type I receptor agonists for treatment of osteoporosis)

RN 862981-03-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5-butyl-7-cyclopentyl- (CA INDEX NAME)



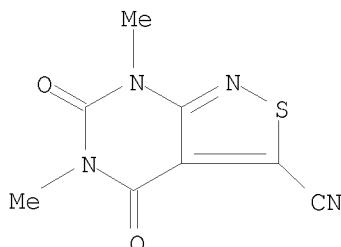
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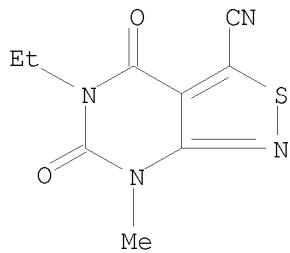
AB Title compds. [I; A, B = S, O, N, CH; R1, R2 = (substituted) alkyl, alkylene, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, cycloalkylaryl, heterocycloaryl], were prepared. Thus, 6-amino-1,3-dibutyl-1H-pyrimidine-2,4-dione in DMF was treated with 1-isothiocyanato-4-methoxybenzene at 23° followed by stirring at 100° for 16 h to give 70% 6-amino-1,3-dibutyl-N-(4-methoxybenzyl)-2,3-dioxo-1,2,3,4-tetrahydropyrimidine-5-carbothioamide. The latter in CHCl₃ was treated with Br₂ followed by stirring for 36 h at 23° to give 34% 3-amino-5,7-dibutylisothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione. I showed binding to PTH1R in membrane prepns. of HEK cells with IC₅₀ = 2-3 μM.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

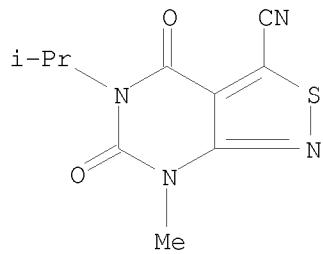
L6 ANSWER 2 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:50046 CAPLUS
 DOCUMENT NUMBER: 138:255198
 TITLE: Novel Synthesis and Reactions of 5,7-Dialkyl-4,6-dioxo-4,5,6,7-tetrahydroisothiazolo[3,4,-d]pyrimidine-3-carbonitriles and 6-Methyl-4-oxo-4H-1-aza-5-oxa-2-thiaindene-3-carbonitrile
 AUTHOR(S): Chang, Yong-Goo; Cho, Hyong Soon; Kim, Kyongtae
 CORPORATE SOURCE: School of Chemistry and Molecular Engineering, Seoul National University, Seoul, 151-742, S. Korea
 SOURCE: Organic Letters (2003), 5(4), 507-510
 PUBLISHER: American Chemical Society
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 138:255198
 IT 502793-19-3P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and substitution of 5,7-dialkyl-4,6-dioxo-4,5,6,7-tetrahydroisothiazolo[3,4,-d]pyrimidine-3-carbonitriles and 6-methyl-4-oxo-4H-1-aza-5-oxa-2-thiaindene-3-carbonitrile)
 RN 502793-19-3 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo- (CA INDEX NAME)



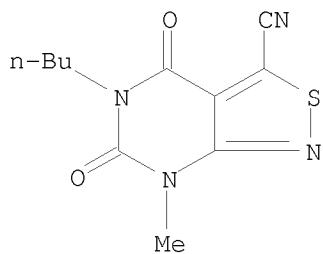
IT 502793-20-6P 502793-21-7P 502793-22-8P
 502793-23-9P 502793-24-0P 502793-25-1P
 502793-26-2P 502793-27-3P 502793-28-4P
 502793-29-5P 502793-30-8P 502793-31-9P
 502793-32-0P 502793-33-1P 502793-34-2P
 502793-35-3P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation and substitution of 5,7-dialkyl-4,6-dioxo-4,5,6,7-tetrahydroisothiazolo[3,4,-d]pyrimidine-3-carbonitriles and 6-methyl-4-oxo-4H-1-aza-5-oxa-2-thiaindene-3-carbonitrile)
 RN 502793-20-6 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 5-ethyl-4,5,6,7-tetrahydro-7-methyl-4,6-dioxo- (CA INDEX NAME)



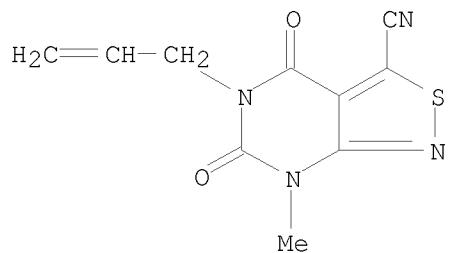
RN 502793-21-7 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-7-methyl-5-(1-methylethyl)-4,6-dioxo- (CA INDEX NAME)



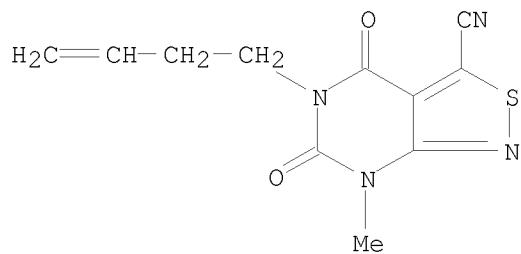
RN 502793-22-8 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 5-butyl-4,5,6,7-tetrahydro-7-methyl-4,6-dioxo- (CA INDEX NAME)



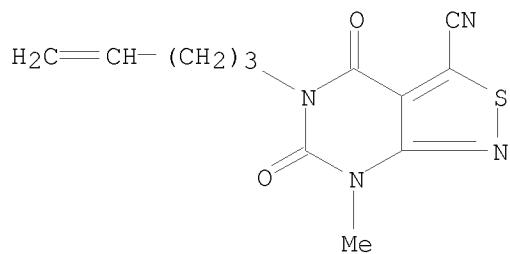
RN 502793-23-9 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-7-methyl-4,6-dioxo-5-(2-propen-1-yl)- (CA INDEX NAME)



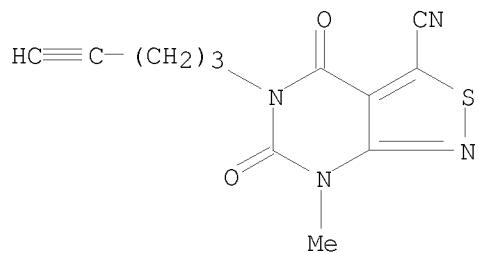
RN 502793-24-0 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 5-(3-buten-1-yl)-4,5,6,7-tetrahydro-7-methyl-4,6-dioxo- (CA INDEX NAME)



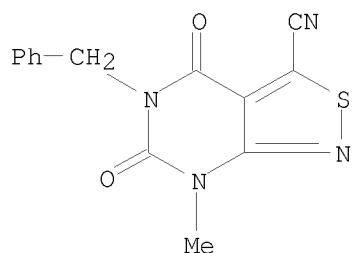
RN 502793-25-1 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-7-methyl-4,6-dioxo-5-(4-penten-1-yl)- (CA INDEX NAME)



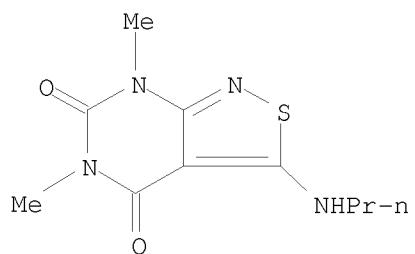
RN 502793-26-2 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-7-methyl-4,6-dioxo-5-(4-pentyne-1-yl)- (CA INDEX NAME)



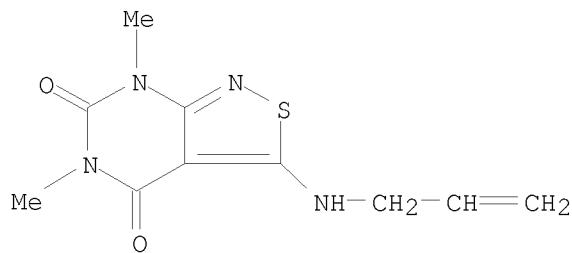
RN 502793-27-3 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-3-carbonitrile, 4,5,6,7-tetrahydro-7-methyl-4,6-dioxo-5-(phenylmethyl)- (CA INDEX NAME)



RN 502793-28-4 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(propylamino)- (CA INDEX NAME)

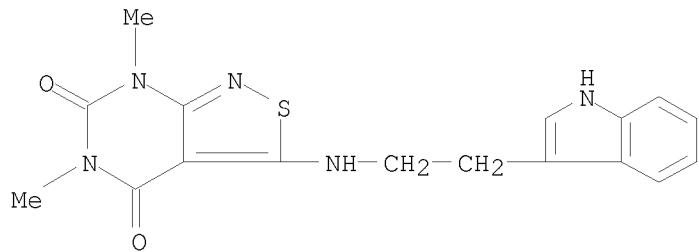


RN 502793-29-5 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(2-propen-1-ylamino)- (CA INDEX NAME)



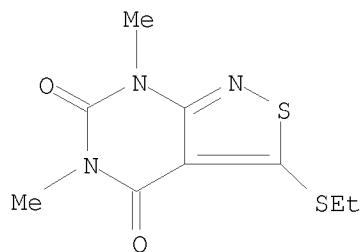
RN 502793-30-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(2-(1H-indol-3-yl)ethyl)amino]-5,7-dimethyl- (CA INDEX NAME)



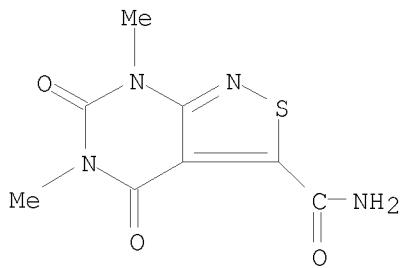
RN 502793-31-9 CAPLUS

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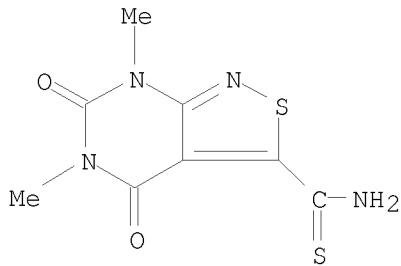
RN 502793-32-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-carboxamide, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo- (CA INDEX NAME)



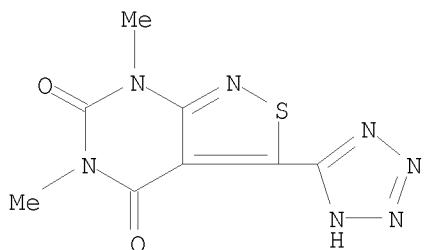
RN 502793-33-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-carbothioamide, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo- (CA INDEX NAME)



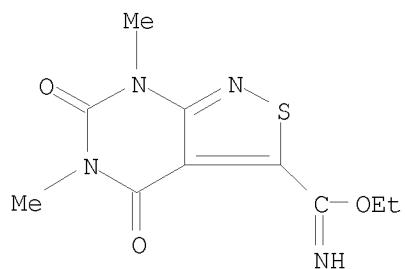
RN 502793-34-2 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(2H-tetrazol-5-yl)- (CA INDEX NAME)



RN 502793-35-3 CAPLUS

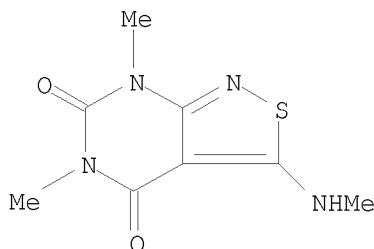
CN Isothiazolo[3,4-d]pyrimidine-3-carboximidic acid, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo-, ethyl ester (CA INDEX NAME)



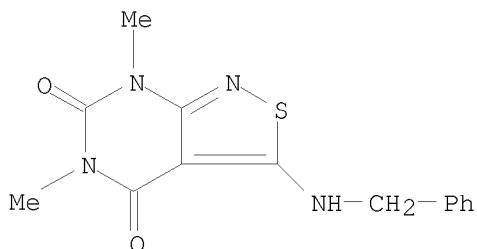
AB 5,7-Dialkyl-4,6-dioxo-4,5,6,7-tetrahydroisothiazolo[3,4-d]pyrimidine-3-carbonitriles, prepared from 6-amino-1,3-dialkyluracils and 4,5-dichloro-5H-1,2,3-dithiazolium chloride (Appel's salt, I), are utilized for the preparation of new derivs. bearing amino, alkylthio, amido, thioamido, tetrazolyl, and carboximidic acid Et ester groups at position 3. Similarly, the reactions of 6-methyl-4-oxo-4H-1-aza-5-oxa-2-thiaindene-3-carbonitrile (II), prepared from 4-amino-6-methyl-2-pyrone and I, with alkyl- and arylamines in DMF at 50 °C and reflux afforded various isothiazole derivs. On the other hand, treatment of II with 1,3-diaminopropane in THF at room temperature, followed by chromatog. on silica gel, gave 3-(2-oxopropyl)-6,7,8-trihydro-4H-1-thia-2,5,9-triazacyclopentacyclononene-4,10-dione in 59% yield.

REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2002:890751 CAPLUS
 DOCUMENT NUMBER: 138:287618
 TITLE: Ring closure reactions of β -nitroso-,
 β -acyl-, and β -thiocarbamoyl- α , β -
 unsaturated sulfilimines. Synthesis of
 [1,2,5]oxadiazolo[3,4-d]-, isoxazolo[3,4-d]-, and
 isothiazolo[3,4-d]pyrimidine derivatives from uracils
 Matsumoto, Nobuaki; Takahashi, Masahiko
 AUTHOR(S):
 CORPORATE SOURCE: Faculty of Engineering, Department of Materials
 Science, Ibaraki University, Hitachi, Ibaraki,
 316-8511, Japan
 SOURCE: Tetrahedron (2002), 58(50), 10073-10079
 CODEN: TETRAB; ISSN: 0040-4020
 PUBLISHER: Elsevier Science Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 138:287618
 IT 61851-90-9P 70425-00-2P 73123-40-7P
 506437-27-0P 506437-28-1P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation and ring closure reactions of β -nitroso-, β -acyl-,
 and β -thiocarbamoyl- α , β -unsatd. sulfilimines)
 RN 61851-90-9 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-
 (methylamino)- (CA INDEX NAME)

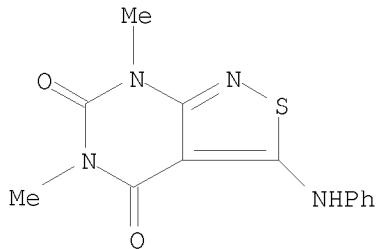


RN 70425-00-2 CAPLUS
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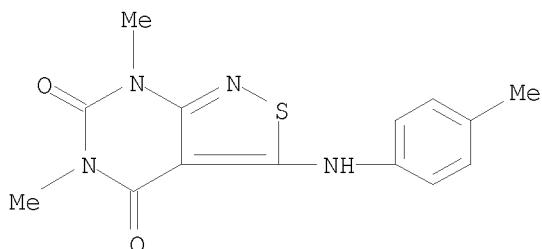
RN 73123-40-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-

(phenylamino)- (CA INDEX NAME)



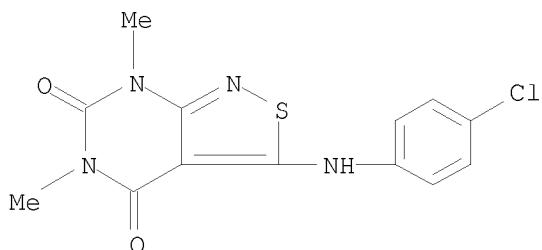
RN 506437-27-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(4-methylphenyl)amino]- (CA INDEX NAME)



RN 506437-28-1 CAPLUS

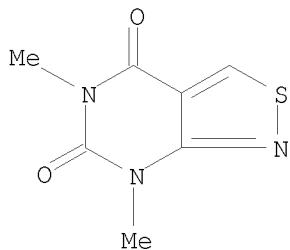
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)amino]-5,7-dimethyl- (CA INDEX NAME)



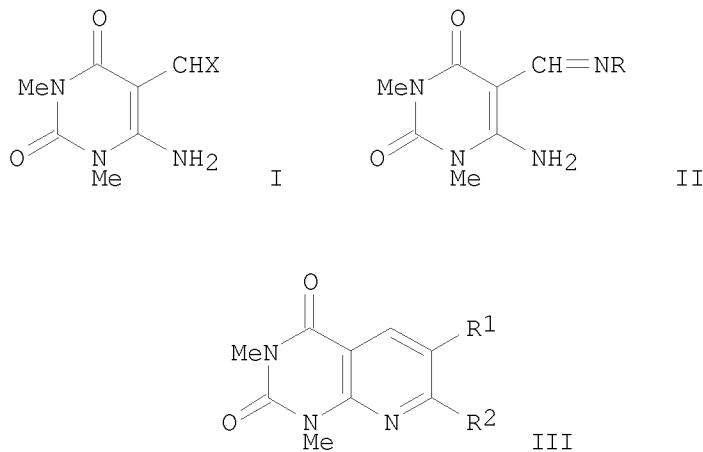
AB 1,3-Dialkyl-6-chlorouracils were treated with S,S-diphenylsulfilimine to give N-(1,3-dialkyluracil-6-yl)-S,S-diphenylsulfilimines. The uracilylsulfilimines were nitrosated, acylated, or thiocarbamoylated to give N-(5-nitroso-, 5-acyl-, or 5-thiocarbamoyluracil-6-yl)sulfilimines, resp. These conjugated sulfilimines were cyclized by thermolysis or photolysis to [1,2,5]oxadiazolo[3,4-d]- isoxazolo[3,4-d]- or isothiazolo[3,4-d]pyrimidine derivs.

REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 4 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1997:213322 CAPLUS
 DOCUMENT NUMBER: 126:293309
 ORIGINAL REFERENCE NO.: 126:56801a,56804a
 TITLE: Reactivities of 6-amino-1,3-dimethyl-5-thioformyluracil toward nucleophiles and its application to synthesis of pyrido[2,3-d]pyrimidines
 Hirota, Kosaku; Kubo, Keiko; Sajiki, Hironao
 AUTHOR(S):
 CORPORATE SOURCE: Laboratory of Medicinal Chemistry, Gifu Pharmaceutical University, Gifu, 502, Japan
 SOURCE: Chemical & Pharmaceutical Bulletin (1997), 45(3), 542-544
 PUBLISHER: Pharmaceutical Society of Japan
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 126:293309
 IT 181465-22-5P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (reactions of thioformyluracil derivative with nucleophiles)
 RN 181465-22-5 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl- (CA INDEX NAME)



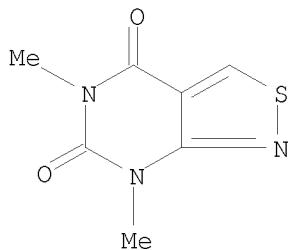
GI



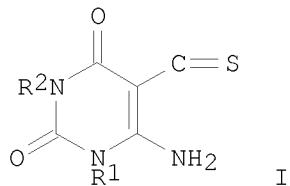
AB The reaction of the 5-thioformyluracil I ($X = S$) with phenylhydrazine and various amines readily afforded the hydrazone II ($R = N\text{HPh}$) and Schiff bases II ($R = \text{OH}$, Me , Ph), resp. Further, carbanions and Wittig reagents reacted with I ($X = S$) to give pyrido[2,3-d]pyrimidines III [$\text{R}1 = \text{cyano}$, CO_2Et , $\text{R}2 = \text{NH}_2$; $\text{R}1 = \text{CO}_2\text{Et}$, $\text{R}2 = \text{OH}$; $\text{R}1 = \text{H}$, $\text{R}2 = \text{NH}_2$]. The corresponding 5-formyluracil I ($X = \text{O}$) possessed much lower reactivities toward these nucleophiles than did I ($X = S$).

REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 5 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1996:457576 CAPLUS
 DOCUMENT NUMBER: 125:221422
 ORIGINAL REFERENCE NO.: 125:41381a, 41384a
 TITLE: Stable thioaldehydes: synthesis, structure assignment, and stability of 6-amino-5-thioformyluracils
 AUTHOR(S): Hirota, Kosaku; Sajiki, Hironao; Kubo, Keiko; Kido, Masaru; Nakagawa, Kazuyuki
 CORPORATE SOURCE: Lab. of Medicinal Chemistry, Gifu Pharmaceutical Univ., Gifu, 502, Japan
 SOURCE: Tetrahedron (1996), 52(30), 9971-9978
 CODEN: TETRAB; ISSN: 0040-4020
 PUBLISHER: Elsevier
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 IT 181465-22-5P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation and properties of (amino)dioxopyrimidinecarbothioaldehydes)
 RN 181465-22-5 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl- (CA INDEX NAME)



GI



AB Stable 6-amino-5-thioformyluracils (i.e., dioxopyrimidinecarbothioaldehydes) I (R1 = Me, Ph, Bu, etc.; R2 = Me, Pr) were synthesized starting from 6-amino-1,3-disubstituted uracils in 23-98% yields. According to the x-ray crystal structure, although I (R1 = Ph, R2 = Me) possessed reasonable double-bond character of the C=S bond, the length of C=S bond in I (R1 = Ph, R2 = Me) is longer than that in of kinetically stabilized thioaldehydes due to the mesomeric effect of the 6-amino group.

L6 ANSWER 6 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1983:107682 CAPLUS

DOCUMENT NUMBER: 98:107682

ORIGINAL REFERENCE NO.: 98:16429a,16432a

TITLE: Glucoside derivatives

PATENT ASSIGNEE(S): Ogura, Haruo, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

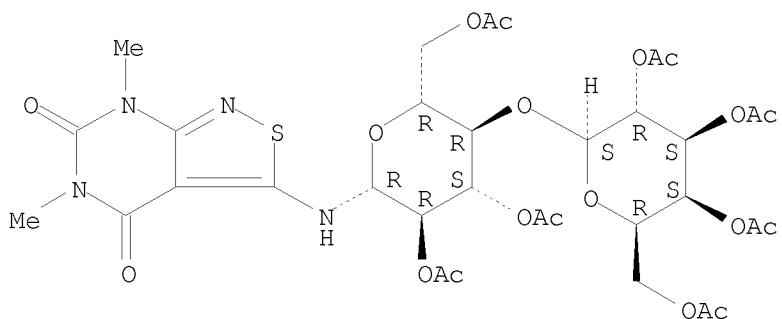
LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

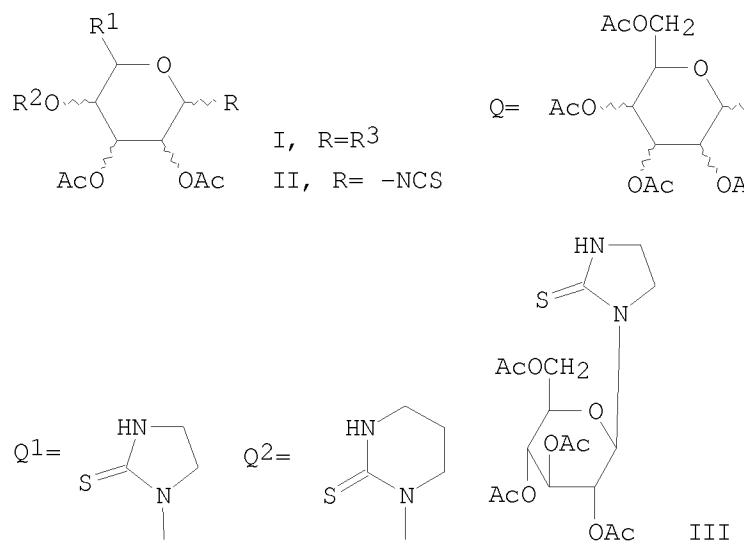
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 57146796	A	19820910	JP 1981-32498	19810309
PRIORITY APPLN. INFO.:			JP 1981-32498	19810309
IT 80681-66-9P				
RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)				
RN 80681-66-9 CAPLUS				
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[2,3,6-tri-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl- β -D-galactopyranosyl)- β -D-glucopyranosyl]amino] - (9CI) (CA INDEX NAME)				

Absolute stereochemistry.



GI



AB Nineteen glycoside derivs. I [$R^1 = H, CH_2OH, CH_2OAc$; $R^2 = Ac, Q$; $R^3 = NHCSNHCH_2CH_2OH, NHCSNHNHCSNH_2, Q_1, Q_2$, etc.] were prepared by reaction of II with the corresponding amino compds. Thus, a mixture of 389 mg β -D-II ($R^1 = CH_2OAc, R^2 = Ac$) and 116 mg $C_1CH_2CH_2NH_2 \cdot HCl$ in pyridine was stirred 3 h at room temperature to give 95% III.

L6 ANSWER 7 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1983:72644 CAPLUS

DOCUMENT NUMBER: 98:72644

ORIGINAL REFERENCE NO.: 98:11139a, 11142a

TITLE: Syntheses of disaccharide isothiocyanates and nucleoside related compounds

AUTHOR(S): Ogura, Haruo; Takahashi, Hiroshi; Kobayashi, Minae

CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, 108, Japan

SOURCE: Nippon Kagaku Kaishi (1982), (10), 1673-81

CODEN: NKAKB8; ISSN: 0369-4577

DOCUMENT TYPE: Journal

LANGUAGE: Japanese

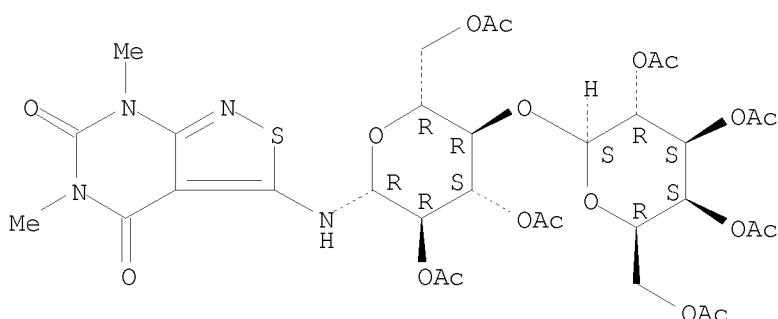
IT 80681-66-9P 84597-66-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 80681-66-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[[2,3,6-tri-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl)- β -D-glucopyranosyl]amino]- (9CI) (CA INDEX NAME)

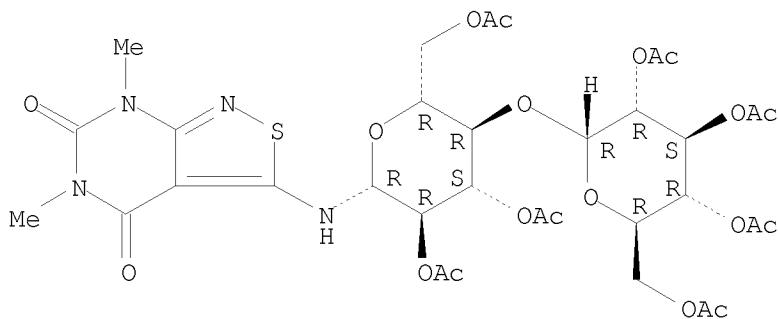
Absolute stereochemistry.



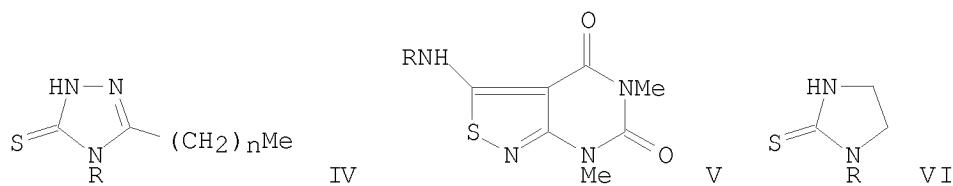
RN 84597-66-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[[2,3,6-tri-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl- α -D-glucopyranosyl)- β -D-glucopyranosyl]amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



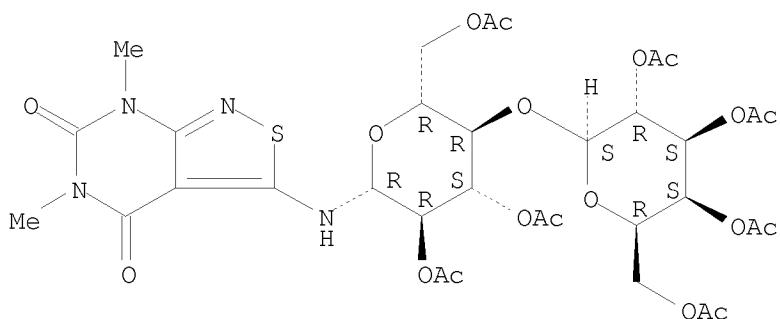
GI



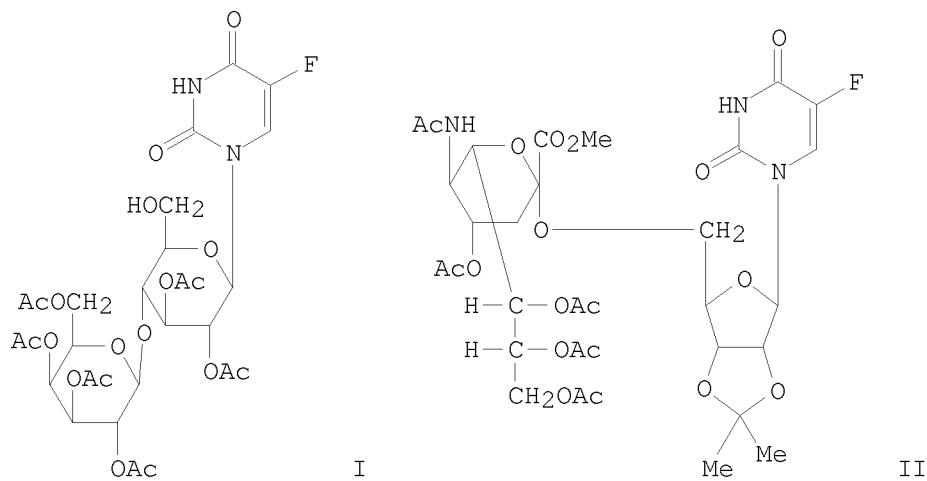
AB Modified nucleoside analogs were prepared starting from hepta-O-acetyl- β -lactosyl isothiocyanate (I), hepta-O-acetyl- β -maltosyl isothiocyanate (II), and hepta-O-acetyl- β -cellobiosyl isothiocyanate (III). I-III reacted with acylhydrazines to give RNHCSNNHCO(CH₂)_nMe (R = disaccharide residue), which afforded triazole disaccharides (IV) by treatment with Ac₂O-H₃PO₄ through cyclodehydration reaction. Reactions of I-III with 6-amino-1,3-dimethyluracil gave disaccharide aminoisothiazolopyrimidines (V) in good yields. Treatment of I-III with 2-amino-2-deoxy- β -D-glucopyranose yielded diglycosylthioureas. Reactions of I-III with chloroethylamine hydrochloride under basic conditions afforded disaccharide imidazolidinethiones (VI) instead of N-glycosyl-N'-chloroethylthioureas.

L6 ANSWER 8 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1982:85908 CAPLUS
 DOCUMENT NUMBER: 96:85908
 ORIGINAL REFERENCE NO.: 96:14119a,14122a
 TITLE: Synthetic O-glycosyl nucleoside analogs
 AUTHOR(S): Ogura, Haruo; Furuhata, Fimio; Iwaki, Kazuo;
 Takahashi, Hiroshi
 CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, 108, Japan
 SOURCE: Nucleic Acids Symposium Series (1981), 10, 23-6
 CODEN: NACSD8; ISSN: 0261-3166
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 IT 80681-66-9P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 80681-66-9 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[2,3,6-tri-
 O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl- β -D-galactopyranosyl)- β -D-
 glucopyranosyl]amino]-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

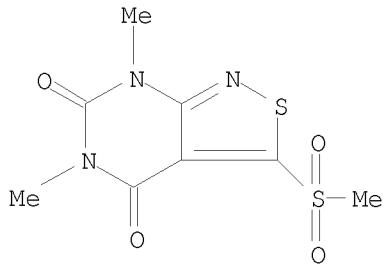


GI

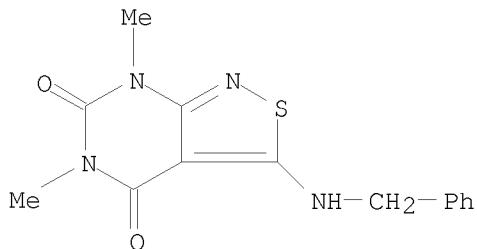


AB O-Glycosyl- and O-neuraminoynucleosides, e.g., I and II, were prepared. I was prepared by condensation of 1,6-anhydro-D-lactose hexaacetate with trimethylsilylated 5-fluorouracil in the presence of SnCl_4 . II was prepared by reaction of Me 2-chloro-4,7,8,9-tetra-O-acetyl-N-acetyl-D-neuraminate with 2',3'-O-isopropylidene-5-fluorouridine.

L6 ANSWER 9 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1980:128845 CAPLUS
 DOCUMENT NUMBER: 92:128845
 ORIGINAL REFERENCE NO.: 92:21015a,21018a
 TITLE: Enamino carbodithioates. III. Methyl
 6-aminouracil-5-carbodithioates. 3. Synthesis and
 reactions of 3-methylthioisothiazolo[3,4-d]-pyrimidine-
 4,6(5H,7H)-diones
 AUTHOR(S): Okuda, Hiroto; Tominaga, Yoshinori; Matsuda, Yoshiro;
 Kobayashi, Goro
 CORPORATE SOURCE: Fac. Pharm. Sci., Nagasaki Univ., Nagasaki, 852, Japan
 SOURCE: Yakugaku Zasshi (1979), 99(10), 989-92
 CODEN: YKKZAJ; ISSN: 0031-6903
 DOCUMENT TYPE: Journal
 LANGUAGE: Japanese
 IT 73123-41-8P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and reactions with amines)
 RN 73123-41-8 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-
 (methylsulfonyl)- (CA INDEX NAME)

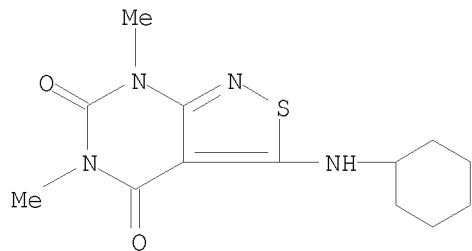


IT 70425-00-2P 70425-05-7P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and substitution reactions of)
 RN 70425-00-2 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-
 [(phenylmethyl)amino]- (CA INDEX NAME)



RN 70425-05-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(cyclohexylamino)-5,7-

dimethyl- (CA INDEX NAME)

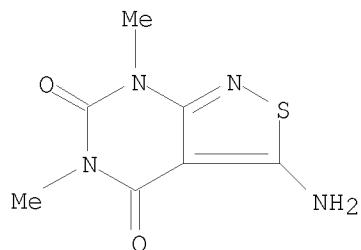


IT 52903-30-7P 52903-33-0P 61851-90-9P
 70425-00-2P 70425-01-3P 70425-02-4P
 70425-03-5P 70425-04-6P 70425-05-7P
 70425-06-8P 70425-07-9P 70425-10-4P
 70425-11-5P 70425-12-6P 70425-13-7P
 73123-40-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

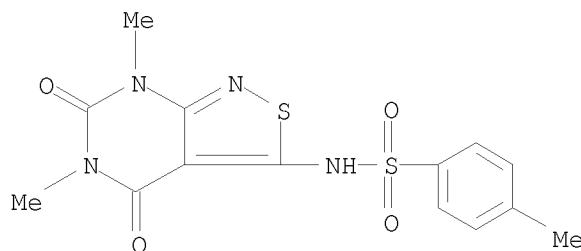
RN 52903-30-7 CAPPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dimethyl- (CA INDEX NAME)



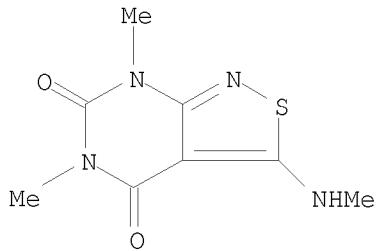
RN 52903-33-0 CAPPLUS

CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



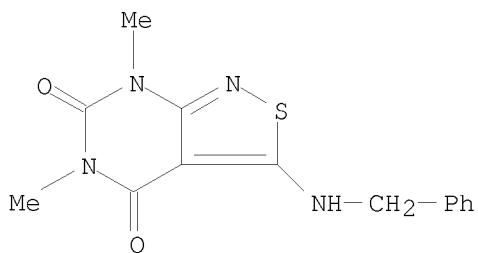
RN 61851-90-9 CAPPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(methylamino)- (CA INDEX NAME)



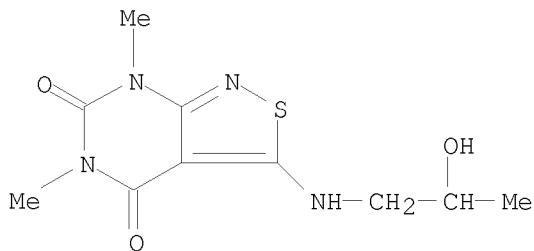
RN 70425-00-2 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(phenylmethyl)amino]- (CA INDEX NAME)



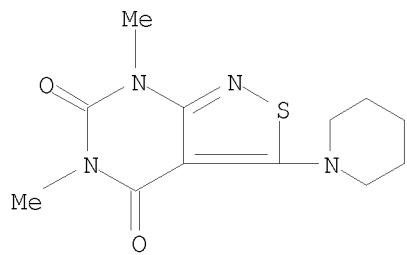
RN 70425-01-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(2-hydroxypropyl)amino]-5,7-dimethyl- (CA INDEX NAME)



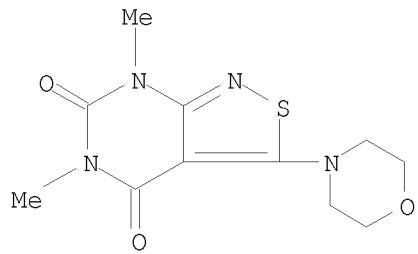
RN 70425-02-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(1-piperidinyl)- (CA INDEX NAME)



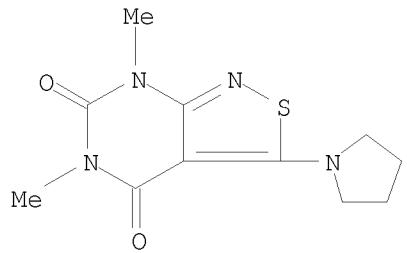
RN 70425-03-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(4-morpholinyl)- (CA INDEX NAME)



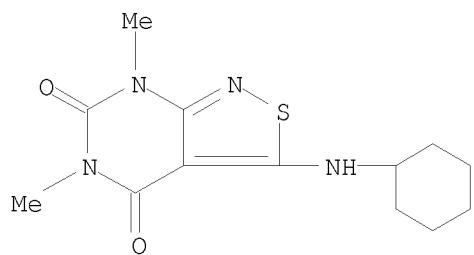
RN 70425-04-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(1-pyrrolidinyl)- (CA INDEX NAME)



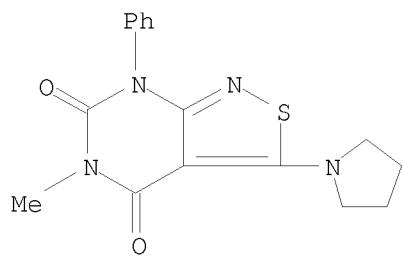
RN 70425-05-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(cyclohexylamino)-5,7-dimethyl- (CA INDEX NAME)



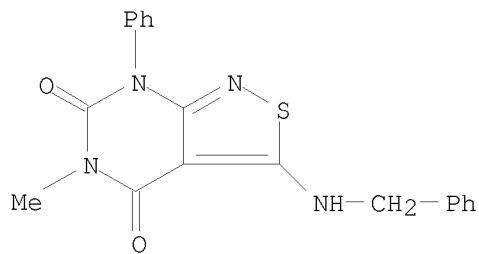
RN 70425-06-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-7-phenyl-3-(1-pyrrolidinyl)- (CA INDEX NAME)



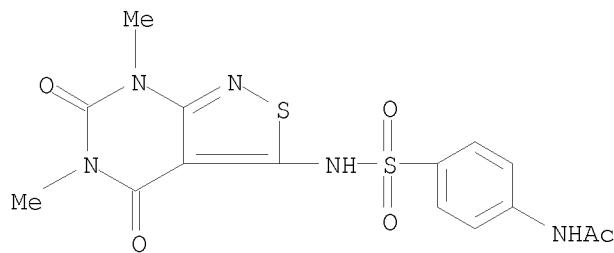
RN 70425-07-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-7-phenyl-3-[(phenylmethyl)amino]- (CA INDEX NAME)



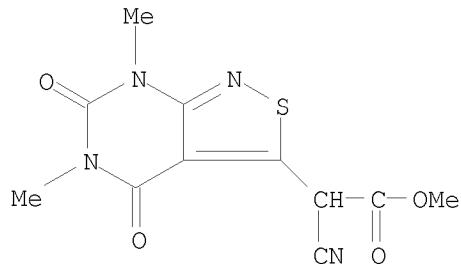
RN 70425-10-4 CAPLUS

CN Acetamide, N-[4-[(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)amino]sulfonyl]phenyl]- (CA INDEX NAME)



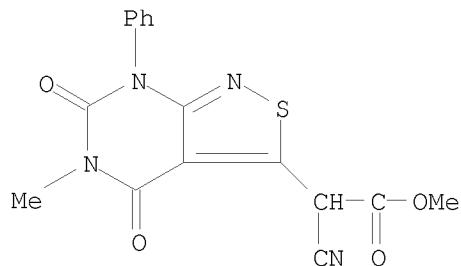
RN 70425-11-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-acetic acid, α -cyano-4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo-, methyl ester (CA INDEX NAME)



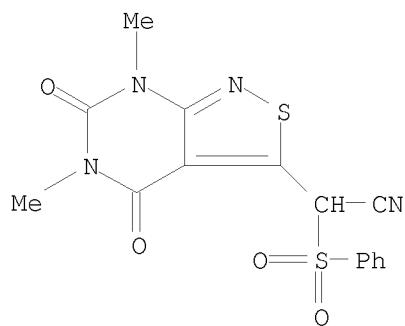
RN 70425-12-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-acetic acid, α -cyano-4,5,6,7-tetrahydro-5-methyl-4,6-dioxo-7-phenyl-, methyl ester (CA INDEX NAME)



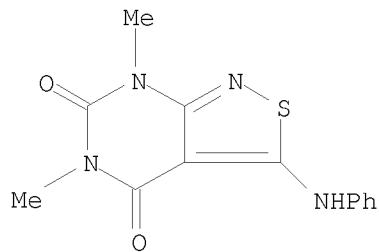
RN 70425-13-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-acetonitrile, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo- α -(phenylsulfonyl)- (CA INDEX NAME)

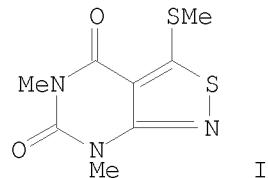


RN 73123-40-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(phenylamino)- (CA INDEX NAME)



GI



AB 3-Methylthioisothiazolopyrimidinediones I (R = Me, Ph) were prepared by treatment with Me 6-aminouracil-5-carbodithioates, prepared by the reaction of 6-aminouracil with CS₂ and Me₂SO₄ in the presence of alkali, with iodine in Me₂SO in a good yield. The reaction of I with amines, amides, and active methylene compds. gave the corresponding substituted products.

L6 ANSWER 10 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1980:22750 CAPLUS

DOCUMENT NUMBER: 92:22750

ORIGINAL REFERENCE NO.: 92:3877a,3880a

TITLE: Studies on heterocyclic compounds. Part XXIX. A one-step synthesis of glycosylaminoisothiazolo[3,4-d]pyrimidines and glycosylaminoisothiazoles

AUTHOR(S): Takahashi, Hiroshi; Nimura, Noriyuki; Ogura, Haruo

CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, 108, Japan

SOURCE: Chemical & Pharmaceutical Bulletin (1979), 27(5), 1147-52

CODEN: CPBTAL; ISSN: 0009-2363

DOCUMENT TYPE: Journal

LANGUAGE: English

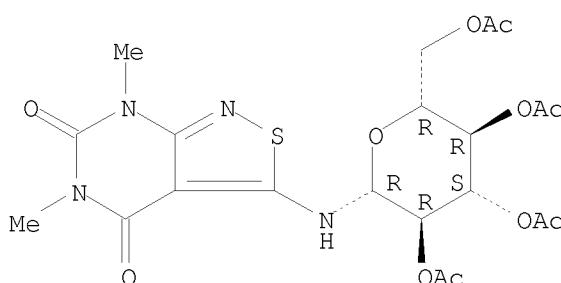
IT 58911-57-2P 62374-42-9P 62374-43-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 58911-57-2 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl)amino]- (9CI) (CA INDEX NAME)

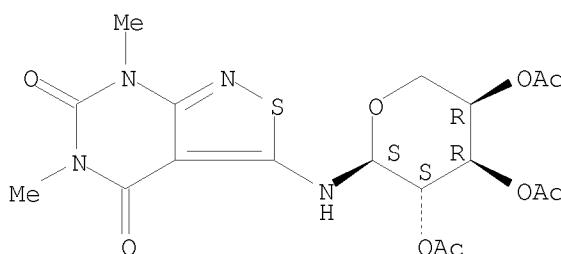
Absolute stereochemistry.



RN 62374-42-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4-tri-O-acetyl- α -D-arabinopyranosyl)amino]- (9CI) (CA INDEX NAME)

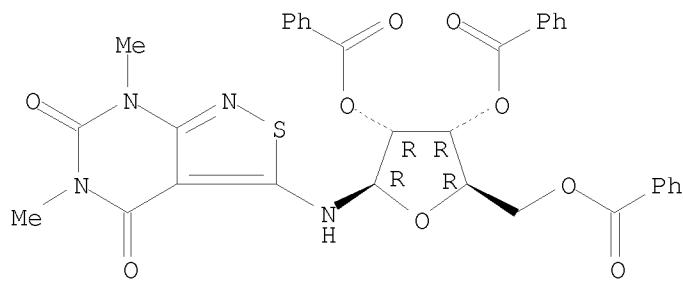
Absolute stereochemistry.



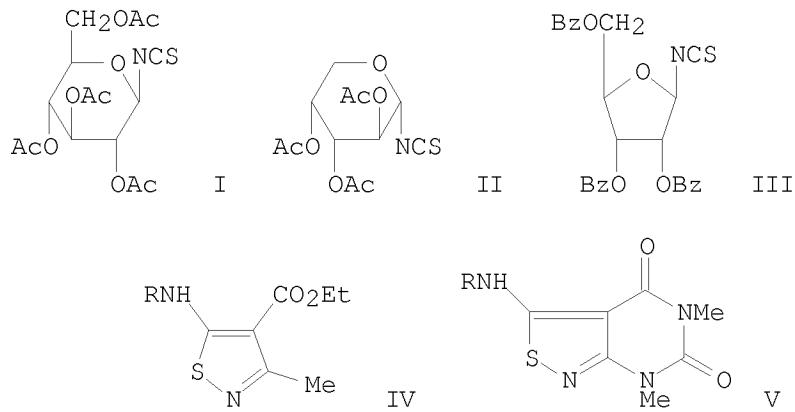
RN 62374-43-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,5-tri-O-benzoyl- β -D-ribofuranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

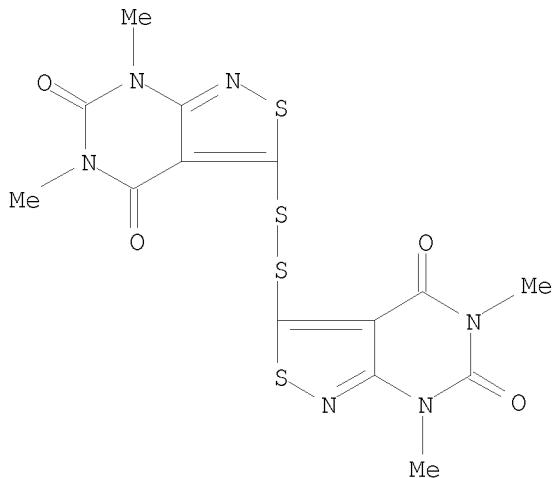


GI

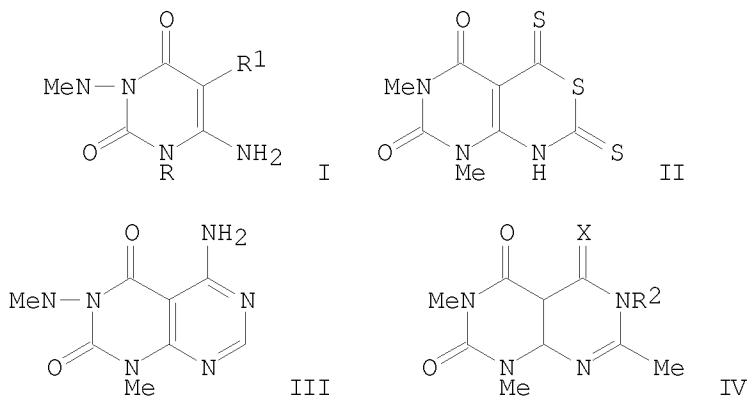


AB The reaction of glycosyl isothiocyanate I with 2-aminopyridine or 2-amino-4-picoline gave N-glycosyl-N'-(2-pyridyl) thiourea and N-glycosyl-N'-(4-methyl-2-pyridyl) thiourea, resp., in good yields; cyclized products were not obtained. On the other hand, the reaction of glycosyl isothiocyanates I, II, and III with $\text{MeC}(\text{NH}_2):\text{CHCO}_2\text{Et}$ gave $\text{MeC}(\text{NH}_2):\text{C}(\text{CSNHR})\text{CO}_2\text{Et}$ ($\text{R} = \text{glycosyl}$) and nucleoside analogs IV ($\text{R} = \text{glycosyl}$). Similar reaction of I-III with 6-amino-1,3-dimethyluracil gave nucleoside analogs V.

L6 ANSWER 11 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1979:575307 CAPLUS
 DOCUMENT NUMBER: 91:175307
 ORIGINAL REFERENCE NO.: 91:28279a,28282a
 TITLE: Enamino carbodithioates. I. Methyl
 6-aminouracil-5-carbodithioates. 1. Synthesis and
 reactions
 AUTHOR(S): Tominaga, Yoshinori; Machida, Tsuyoshi; Okuda, Hiroto;
 Matsuda, Yoshiro; Kobayashi, Goro
 CORPORATE SOURCE: Fac. Pharm. Sci., Nagasaki Univ., Nagasaki, 852, Japan
 SOURCE: Yakugaku Zasshi (1979), 99(5), 515-20
 CODEN: YKKZAJ; ISSN: 0031-6903
 DOCUMENT TYPE: Journal
 LANGUAGE: Japanese
 OTHER SOURCE(S): CASREACT 91:175307
 IT 71266-55-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 71266-55-2 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3,3'-dithiobis[5,7-dimethyl-
 (9CI) (CA INDEX NAME)



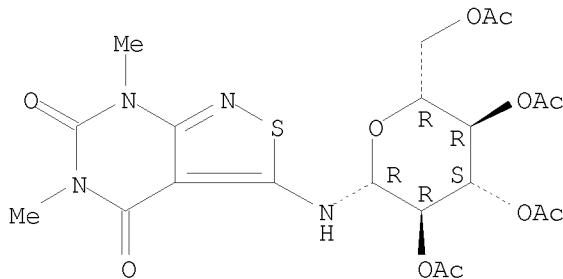
GI



AB The 6-aminouracils I ($R = Me, Ph, R1 = H$) treated with CS₂ and Me₂SO₄ in NaOH-DMSO gave dithiocarboxylates I ($R = MePh; R1 = CS_2Me$). An excess of CS₂ in the previous reaction with I ($R = Me, R1 = H$) gave the pyrimido[4,5-d][1,3]thiazine II in good yield. The reaction of I ($R = Me, Ph; R1 = CS_2Me$) and R₂NH ($R2 = H, NH_2, Me, PhCH_2, cyclohexyl, morpholino, HOCH_2CH_2, (EtO)_2CHCH_2$), yielded thiocarbamoyluracils I [$R = Me, Ph, R1 = C(S)NR_2$] in 31-91% yields. I [$R = Me, R1 = C(S)SMe$] and HCONH₂ reacted to yield III. Also prepared were pyrimido[4,5-d]pyrimidines IV ($R2 = H, Me; X = S$) and IV ($R2 = PhCH_2, X = O$) from the reaction of I [$R = Me, R1 = C(S)NR_2, R2 = H, Me, PhCH_2$] with Ac₂O.

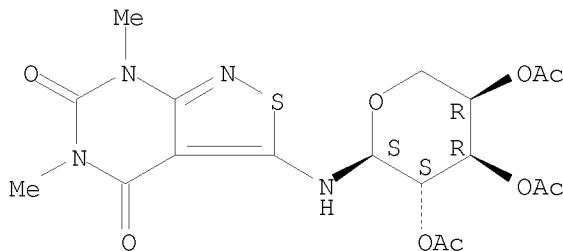
L6 ANSWER 12 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1979:439764 CAPLUS
 DOCUMENT NUMBER: 91:39764
 ORIGINAL REFERENCE NO.: 91:6497a,6500a
 TITLE: Syntheses of nucleoside analogs using glycosyl isothiocyanate
 AUTHOR(S): Ogura, Haruo; Takahashi, Hiroshi
 CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, Japan
 SOURCE: Tennen Yuki Kagobutsu Toronkai Koen Yoshishu, 21st (1978), 221-8. Hokkaido Daigaku Nogakubu: Sapporo, Japan.
 CODEN: 39NQAF
 DOCUMENT TYPE: Conference
 LANGUAGE: Japanese
 IT 58911-57-2P 62374-42-9P 62374-43-0P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 58911-57-2 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



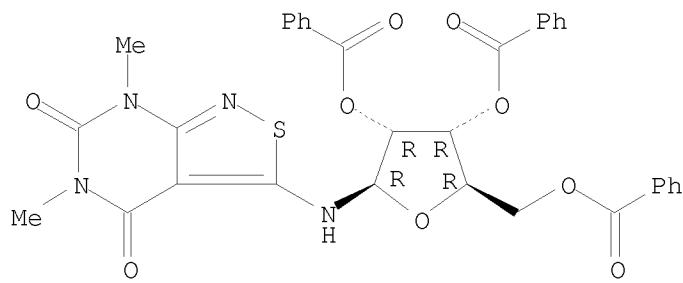
RN 62374-42-9 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4-tri-O-acetyl- α -D-arabinopyranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

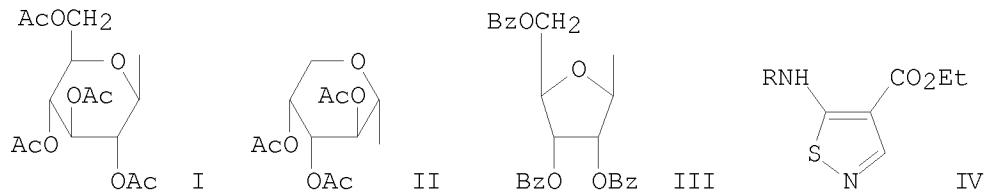


RN 62374-43-0 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,5-tri-O-benzoyl- β -D-ribofuranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



GI



AB Various nucleoside analogs containing isothiazole, isothiazolopyrimidine, fused imidazole, pyrimidothiadiazine, pyrazolopyrimidine, triazole, or triazine moieties were prepared by using RNCS ($R = I$, II , or III). E.g., reaction of RNCS and $\text{MeC}(\text{NH}_2):\text{CHCO}_2\text{Et}$ gave (glycosylamino)isothiazoles IV and $\text{MeC}(\text{NH}_2):\text{C}(\text{SCNHR})\text{CO}_2\text{Et}$ (V). V readily cyclized to IV .

L6 ANSWER 13 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1979:405191 CAPLUS

DOCUMENT NUMBER: 91:5191

ORIGINAL REFERENCE NO.: 91:975a,978a

TITLE: Enamino dithiocarboxylates. II. Methyl 6-aminouracil-5-dithiocarboxylates. 2. Synthesis and reactions of 3-methylthioisothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-diones

AUTHOR(S): Okuda, Hiroto; Tominaga, Yoshinori; Matsuda, Yoshiro; Kobayashi, Goro

CORPORATE SOURCE: Fac. Pharm. Sci., Nagasaki Univ., Nagasaki, 852, Japan

SOURCE: Heterocycles (1979), 12(4), 485-8

CODEN: HTCYAM; ISSN: 0385-5414

DOCUMENT TYPE: Journal

LANGUAGE: English

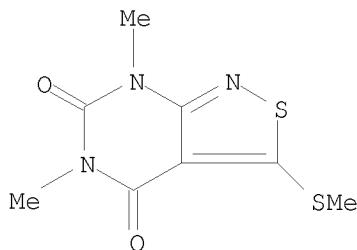
OTHER SOURCE(S): CASREACT 91:5191

IT 70424-98-5P 70424-99-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and nucleophilic substitutions of)

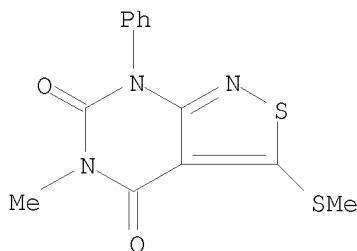
RN 70424-98-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(methylthio)-(CA INDEX NAME)



RN 70424-99-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-3-(methylthio)-7-phenyl- (CA INDEX NAME)



IT 52903-33-0P 61851-90-9P 70425-00-2P

70425-01-3P 70425-02-4P 70425-03-5P

70425-04-6P 70425-05-7P 70425-06-8P

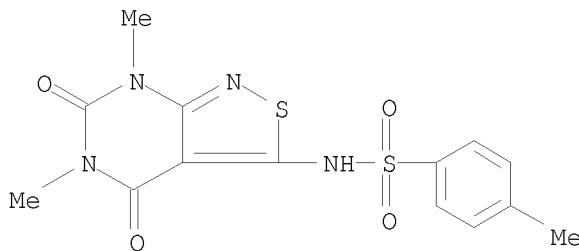
70425-07-9P 70425-10-4P 70425-11-5P

70425-12-6P 70425-13-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

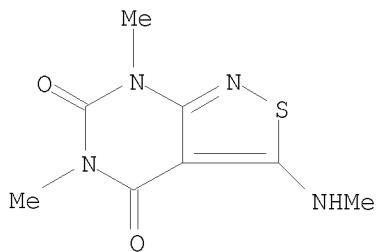
RN 52903-33-0 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



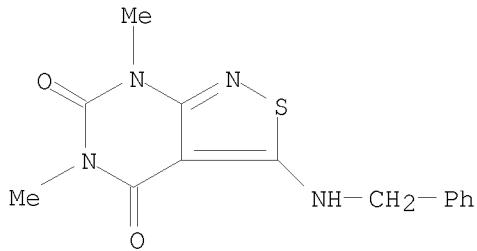
RN 61851-90-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(methylamino)- (CA INDEX NAME)



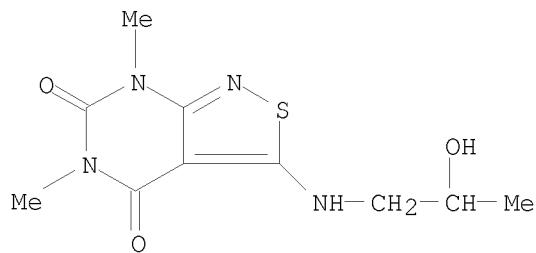
RN 70425-00-2 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(phenylmethyl)amino]- (CA INDEX NAME)



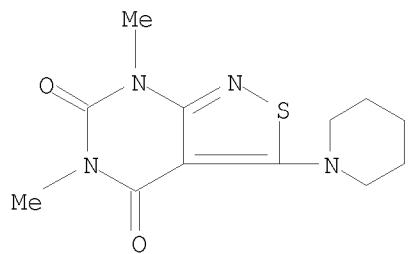
RN 70425-01-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(2-hydroxypropyl)amino]-5,7-dimethyl- (CA INDEX NAME)



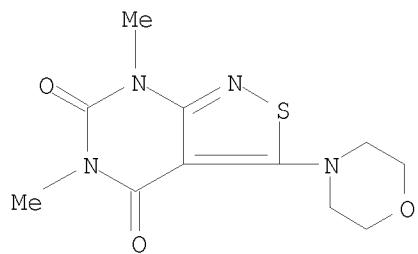
RN 70425-02-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(1-piperidinyl)- (CA INDEX NAME)



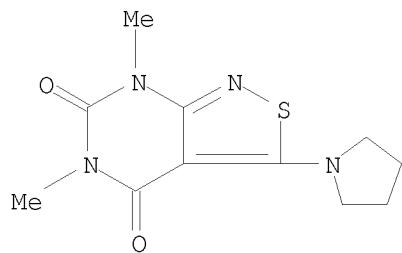
RN 70425-03-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(4-morpholinyl)- (CA INDEX NAME)

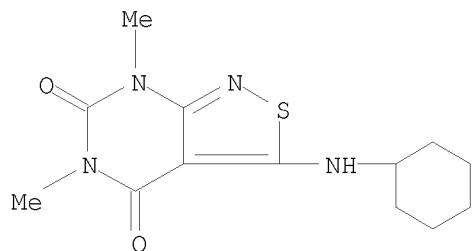


RN 70425-04-6 CAPLUS

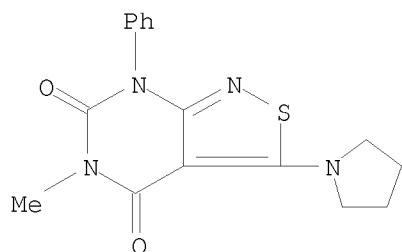
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(1-pyrrolidinyl)- (CA INDEX NAME)



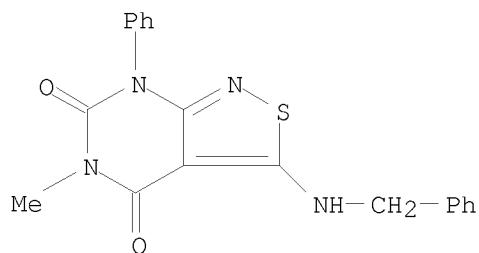
RN 70425-05-7 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(cyclohexylamino)-5,7-dimethyl- (CA INDEX NAME)



RN 70425-06-8 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-7-phenyl-3-(1-pyrrolidinyl)- (CA INDEX NAME)

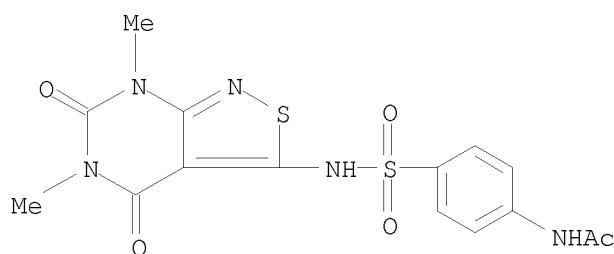


RN 70425-07-9 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-7-phenyl-3-[(phenylmethyl)amino]- (CA INDEX NAME)



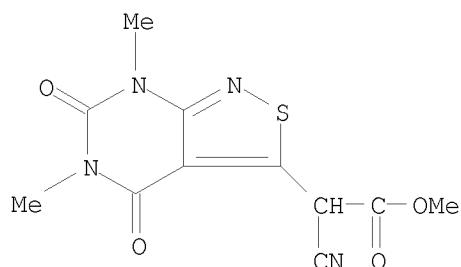
RN 70425-10-4 CAPLUS

CN Acetamide, N-[4-[(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)amino]sulfonyl]phenyl- (CA INDEX NAME)



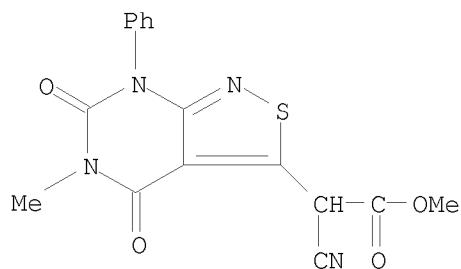
RN 70425-11-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-3-acetic acid, α -cyano-4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo-, methyl ester (CA INDEX NAME)

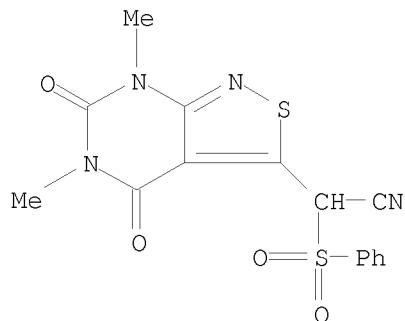


RN 70425-12-6 CAPLUS

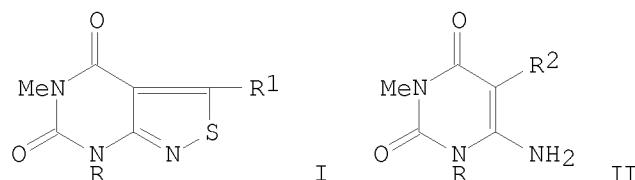
CN Isothiazolo[3,4-d]pyrimidine-3-acetic acid, α -cyano-4,5,6,7-tetrahydro-5-methyl-4,6-dioxo-7-phenyl-, methyl ester (CA INDEX NAME)



RN 70425-13-7 CAPLUS

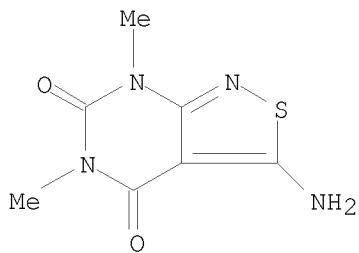
CN Isothiazolo[3,4-d]pyrimidine-3-acetonitrile, 4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxo- α -(phenylsulfonyl)- (CA INDEX NAME)

GI



AB Isothiazolopyrimidinediones I ($R = Me, Ph$; $R1 = SMe$) were obtained by treating uracils II ($R2 = H$) with CS_2 and Me_2SO_4 and cyclizing II ($R2 = CS_2Me$) with iodine. I ($R1 = SMe$) underwent nucleophilic substitution to give I [$R1 = amino, CH(CN)CO_2Me, CH(CN)SO_2Ph$].

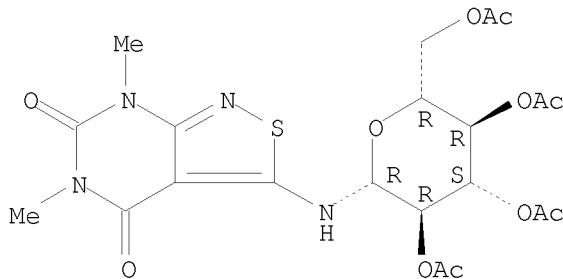
L6 ANSWER 14 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1978:580279 CAPLUS
 DOCUMENT NUMBER: 89:180279
 ORIGINAL REFERENCE NO.: 89:28019a,28022a
 TITLE: Syntheses of modified nucleoside analogs with
 isothiocyanates
 AUTHOR(S): Ogura, Haruo; Takahashi, Hiroshi
 CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, Japan
 SOURCE: Symp. Heterocycl., [Pap.] (1977), 135-40. Editor(s):
 Kametani, Tetsuji. Sendai Inst. Heterocycl. Chem.:
 Sendai, Japan.
 CODEN: 38WUAV
 DOCUMENT TYPE: Conference
 LANGUAGE: English
 IT 52903-30-7DP, glycosyl derivs.
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 52903-30-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dimethyl- (CA
 INDEX NAME)



AB Syntheses of nucleoside analogs via reactions of glycosyl isothiocyanate and glyconyl isothiocyanate with amino acids, enamines, and diamines are outlined.

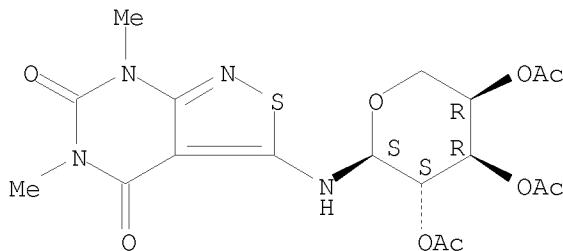
L6 ANSWER 15 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1977:140380 CAPLUS
 DOCUMENT NUMBER: 86:140380
 ORIGINAL REFERENCE NO.: 86:22061a,22064a
 TITLE: Reaction of enamine with D-glycosyl and D-gluconyl isothiocyanate
 AUTHOR(S): Ogura, Haruo; Takahashi, Hiroshi; Takeda, Kazuyoshi; Nimura, Noriyuki
 CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, Japan
 SOURCE: Nucleic Acids Research, Special Publication (1976), 2 (Symp. Nucleic Acids Chem., 4th, 1976), 7-10
 CODEN: NARPD6; ISSN: 0309-1872
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 IT 58911-57-2P 62374-42-9P 62374-43-0P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 58911-57-2 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



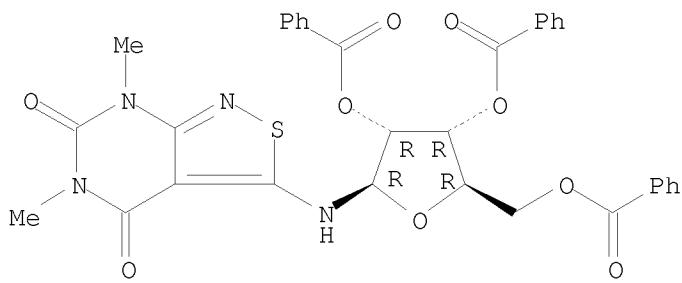
RN 62374-42-9 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4-tri-O-acetyl- α -D-arabinopyranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

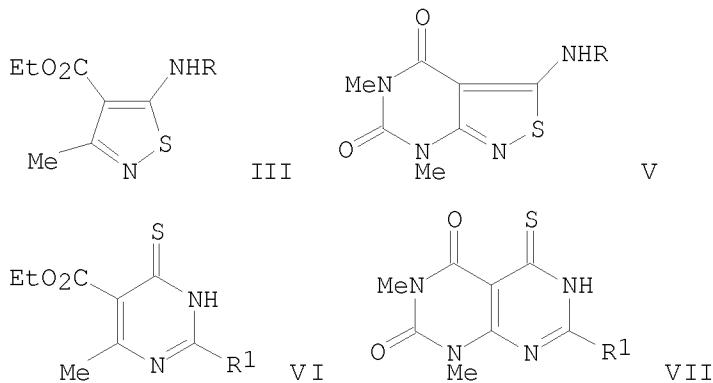


RN 62374-43-0 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,5-tri-O-benzoyl- β -D-ribofuranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



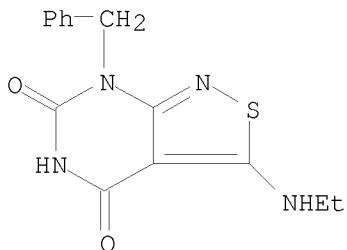
GI



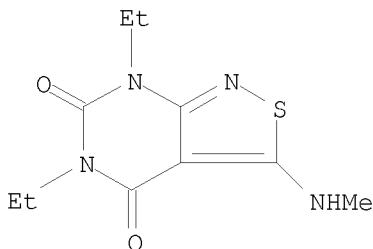
AB Treatment of RNCS (I; R = tetra-O-acetyl- β -D-glucopyranosyl, tri-O-acetyl- α -D-ribopyranosyl, tri-O-acetyl- α -D-arabinopyranosyl, tri-O-benzoyl- β -D-ribofuranosyl) with H₂NCMe:CHCO₂Et (II) in THF at room temperature gave isothiazoles III and H₂NCMe:C(CO₂Et)C(S)NHR, and I cyclized with 6-amino-1,3-dimethyluracil (IV) to give thiazolopyrimidines V. Treatment of penta-O-acetyl-D-gluconyl isothiocyanate with II and IV gave VI and VII (R₁ = penta-O-acetyl-D-gluconyl).

L6 ANSWER 16 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1977:89864 CAPLUS
 DOCUMENT NUMBER: 86:89864
 ORIGINAL REFERENCE NO.: 86:14197a,14200a
 TITLE: Isothiazolo[3,4-d]pyrimidines
 INVENTOR(S): Furukawa, Yoshiyasu; Shima, Shunsuke
 PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 11 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 51091294	A	19760810	JP 1975-16658	19750207
PRIORITY APPLN. INFO.:				
IT 60663-75-4P	60663-76-5P	60663-78-7P	JP 1975-16658	A 19750207
60663-79-8P	60663-80-1P	60663-83-4P		
60663-84-5P	61851-90-9P			
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation and alkylation of)				
RN 60663-75-4	CAPLUS			
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylamino)-7-(phenylmethyl)-	(CA INDEX NAME)			

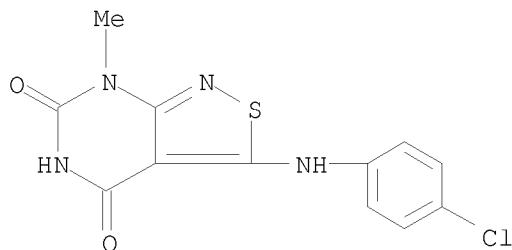


RN 60663-76-5 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-(methylamino)-
 (CA INDEX NAME)



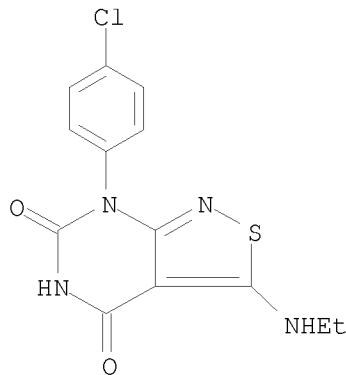
RN 60663-78-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)amino]-7-methyl- (CA INDEX NAME)



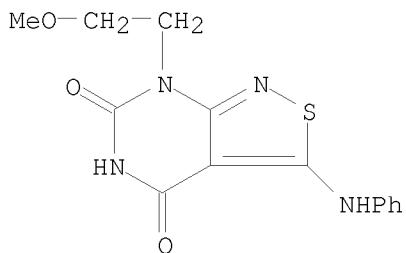
RN 60663-79-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-3-(ethylamino)- (CA INDEX NAME)



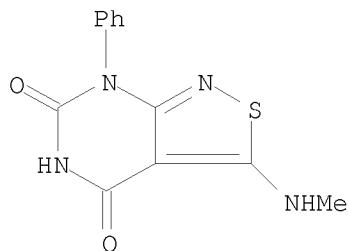
RN 60663-80-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methoxyethyl)-3-(phenylamino)- (CA INDEX NAME)



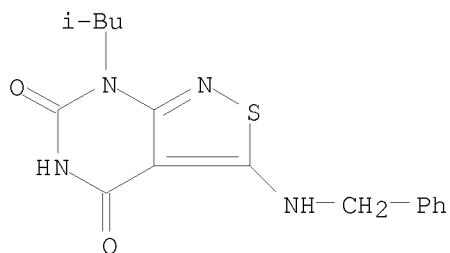
RN 60663-83-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(methylamino)-7-phenyl- (CA INDEX NAME)



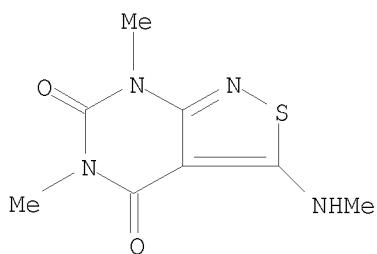
RN 60663-84-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methylpropyl)-3-[(phenylmethyl)amino]- (CA INDEX NAME)



RN 61851-90-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-(methylamino)- (CA INDEX NAME)



IT 56540-86-4P 56540-98-8P 56541-01-6P

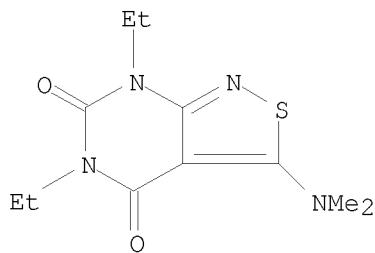
56541-02-7P 56541-03-8P 56541-04-9P

56541-05-0P 60663-90-3P 61851-88-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

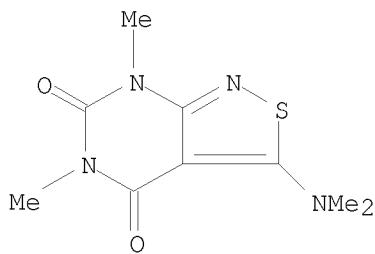
RN 56540-86-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-diethyl- (CA INDEX NAME)



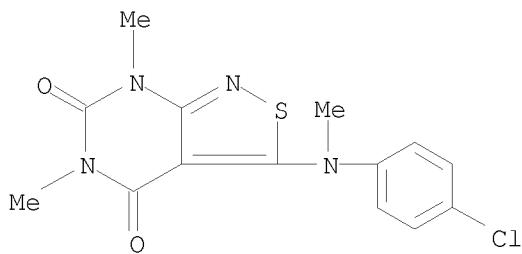
RN 56540-98-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-dimethyl- (CA INDEX NAME)



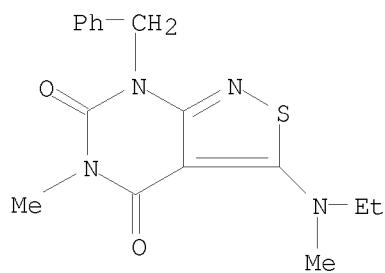
RN 56541-01-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)methylamino]-5,7-dimethyl- (CA INDEX NAME)

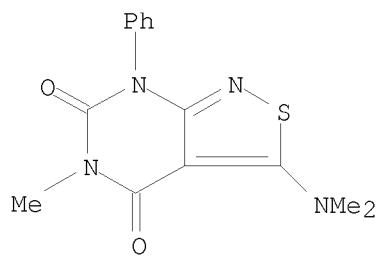


RN 56541-02-7 CAPLUS

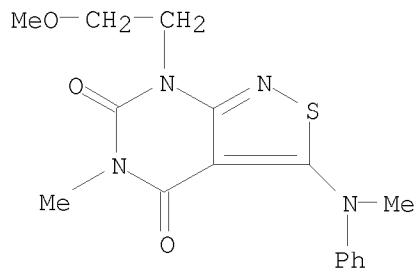
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylmethylamino)-5-methyl-7-(phenylmethyl)- (CA INDEX NAME)



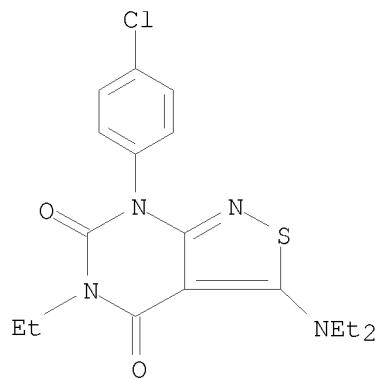
RN 56541-03-8 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5-methyl-7-phenyl- (CA INDEX NAME)



RN 56541-04-9 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methoxyethyl)-5-methyl-3-(methylphenylamino)- (CA INDEX NAME)

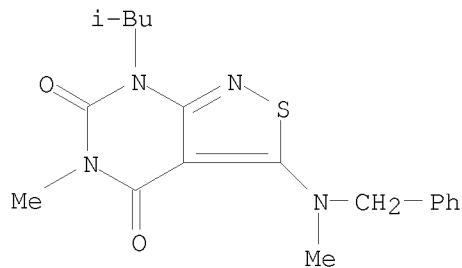


RN 56541-05-0 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-3-(diethylamino)-5-ethyl- (CA INDEX NAME)



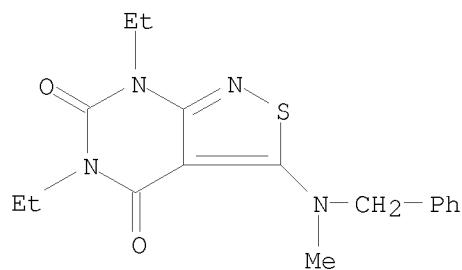
RN 60663-90-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-3-[methyl(phenylmethyl)amino]-7-(2-methylpropyl)- (CA INDEX NAME)

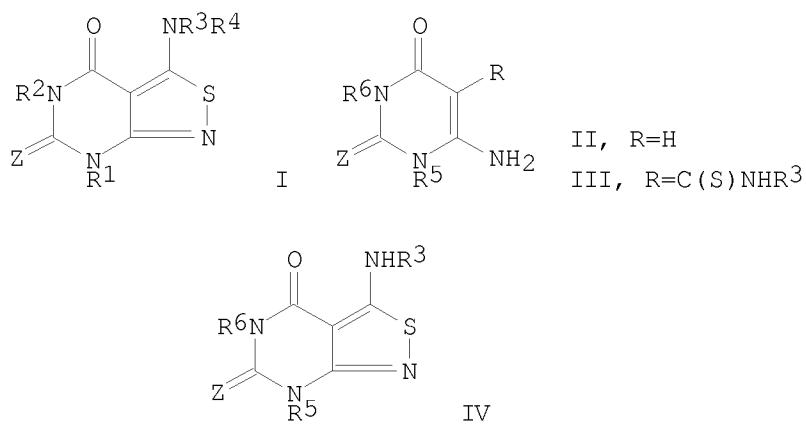


RN 61851-88-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,6H)-dione, 5,7-diethyl-3-[methyl(phenylmethyl)amino]- (9CI) (CA INDEX NAME)

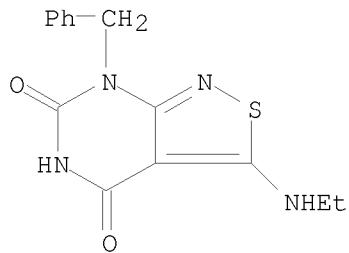


GI

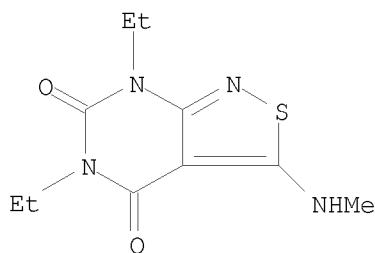


AB Isothiazolo[3,4-d]pyrimidines I ($R_1, R_3 = \text{alkyl, aryl}; R_2, R_4 = \text{alkyl}; Z = O, S$) were prepared by reaction of II ($R_5, R_6 = H, \text{alkyl, aryl}$) with $R_3\text{NCS}$, cyclization of the resulting III, and alkylation of the resulting IV. I had adenosine 3',5'-cyclic phosphate phosphodiesterase-inhibiting activity and are useful as antiinflammatory agents and sedatives (no data). Thus, 7.32 g II ($R_5 = R_6 = \text{Et}, Z = O$) in pyridine was refluxed with 8 ml MeNCS 24 h to give 6.74 g III ($R_3 = \text{Me}, R_5 = R_6 = \text{Et}, Z = O$) (V). Stirring 5.8 g V in AcOEt with 3.4 ml Br 1 h at room temperature gave 5 g IV ($R_3 = \text{Me}, R_5 = R_6 = \text{Et}, Z = O$) (VI). Stirring a mixture of 5 g VI, 5 g K_2CO_3 , and 5 ml MeI in DMF 16 h at room temperature gave 4 g I ($R_1 = R_2 = \text{Et}, R_3 = R_4 = \text{Me}, Z = O$). I also prepared were (R_1, R_2, R_3, R_4, Z given): Me, Me, Me, Me, O; Ph, Me, Me, Me, O; $p\text{-ClC}_6\text{H}_4$, Et, Et, Et, O; Me, Me, $p\text{-ClC}_6\text{H}_4$, Me, O; PhCH_2 , Me, Et, Me, O; $\text{MeOCH}_2\text{CH}_2$, Me, Ph, Me, O; Et, Et, Me, Me, S; iso-Bu, Me, PhCH_2 , Me, O; and Et, Et, PhCH_2 , Me, O.

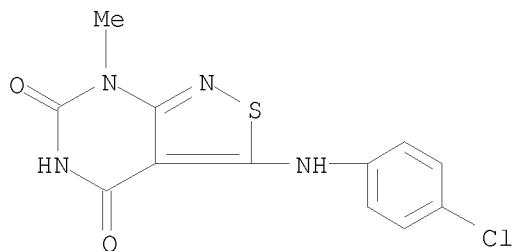
L6 ANSWER 17 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1976:560012 CAPLUS
 DOCUMENT NUMBER: 85:160012
 ORIGINAL REFERENCE NO.: 85:25613a,25616a
 TITLE: Fused pyrimidines. II. Synthesis and oxidation of
 3-aminoisothiazolo[3,4-d]pyrimidines
 AUTHOR(S): Furukawa, Yoshiyasu; Shima, Shunsuke
 CORPORATE SOURCE: Cent. Res. Div., Takeda Chem. Ind., Ltd., Osaka, Japan
 SOURCE: Chemical & Pharmaceutical Bulletin (1976), 24(5),
 979-86
 CODEN: CPBTAL; ISSN: 0009-2363
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 85:160012
 IT 60663-75-4P 60663-76-5P 60663-78-7P
 60663-79-8P 60663-83-4P 60663-84-5P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and alkylation of)
 RN 60663-75-4 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylamino)-7-
 (phenylmethyl)- (CA INDEX NAME)



RN 60663-76-5 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-(methylamino)-
 (CA INDEX NAME)

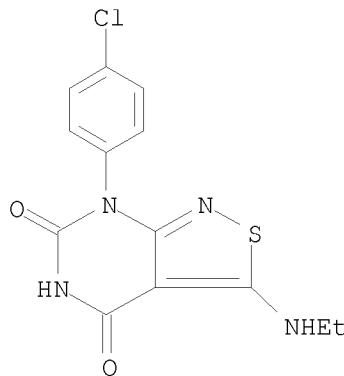


RN 60663-78-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)amino]-7-
 methyl- (CA INDEX NAME)



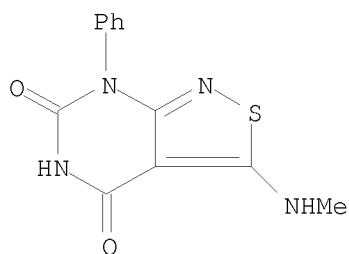
RN 60663-79-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-3-(ethylamino)- (CA INDEX NAME)



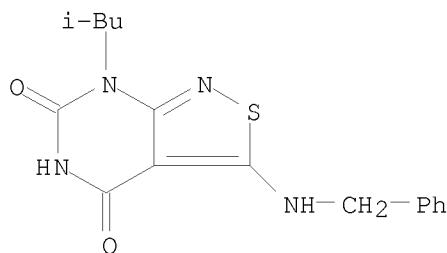
RN 60663-83-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(methylamino)-7-phenyl- (CA INDEX NAME)



RN 60663-84-5 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methylpropyl)-3-[(phenylmethyl)amino]- (CA INDEX NAME)

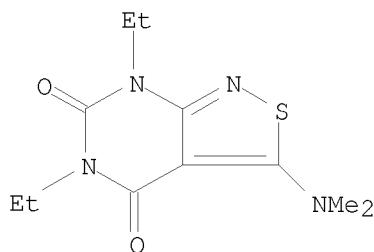


IT 56540-86-4P 56541-01-6P 56541-02-7P
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 60663-77-6P 60663-80-1P 60663-81-2P
 60663-82-3P 60663-85-6P 60663-86-7P
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 60663-90-3P 60700-23-4P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

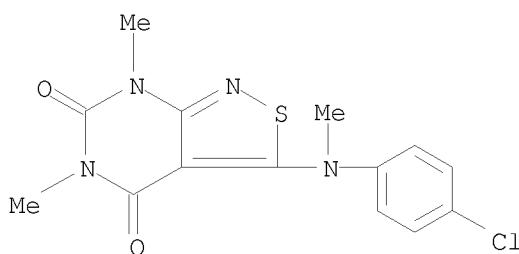
RN 56540-86-4 CAPPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-diethyl- (CA INDEX NAME)



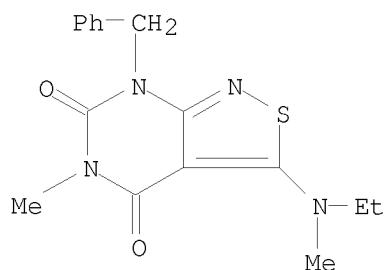
RN 56541-01-6 CAPPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)methylamino]-5,7-dimethyl- (CA INDEX NAME)



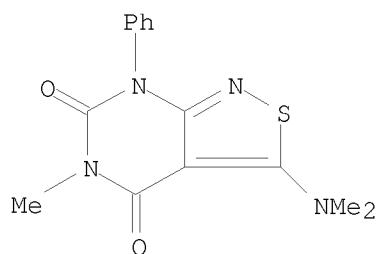
RN 56541-02-7 CAPPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylmethylamino)-5-methyl-7-(phenylmethyl)- (CA INDEX NAME)



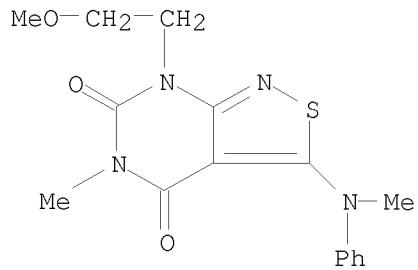
RN 56541-03-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5-methyl-7-phenyl- (CA INDEX NAME)



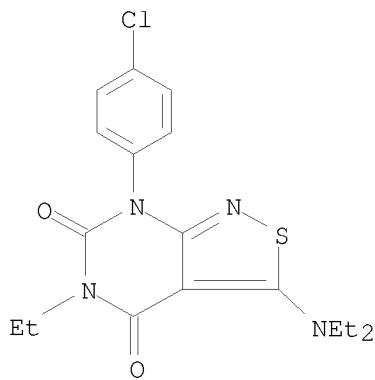
RN 56541-04-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methoxyethyl)-5-methyl-3-(methylphenylamino)- (CA INDEX NAME)



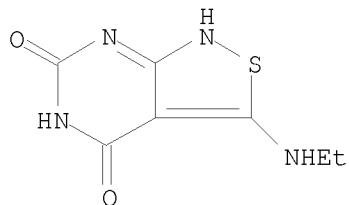
RN 56541-05-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-3-(diethylamino)-5-ethyl- (CA INDEX NAME)



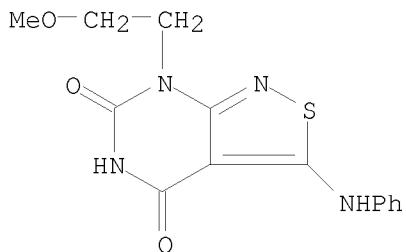
RN 60663-77-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylamino)- (9CI) (CA INDEX NAME)



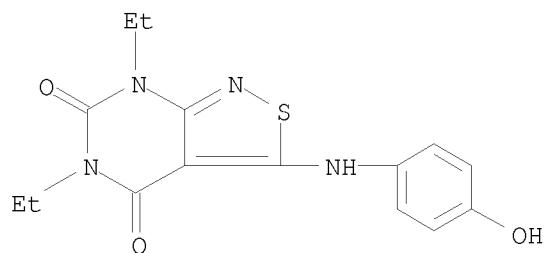
RN 60663-80-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methoxyethyl)-3-(phenylamino)- (CA INDEX NAME)



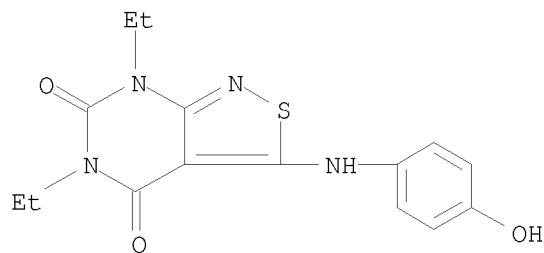
RN 60663-81-2 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-[(4-hydroxyphenyl)amino]- (CA INDEX NAME)



RN 60663-82-3 CAPLUS

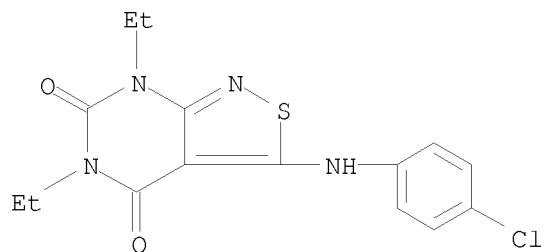
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-[(4-hydroxyphenyl)amino]-, dihydروبromide (9CI) (CA INDEX NAME)



● 2 HBr

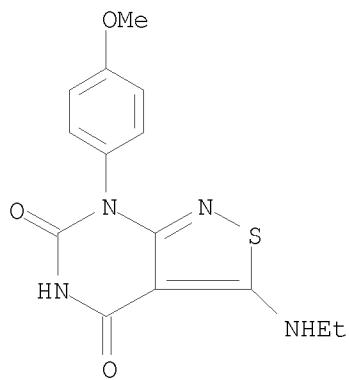
RN 60663-85-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)amino]-5,7-diethyl- (CA INDEX NAME)



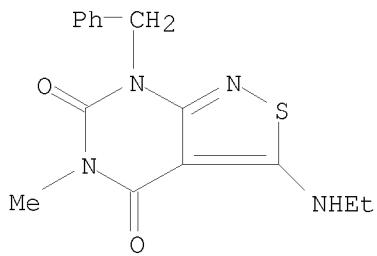
RN 60663-86-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylamino)-7-(4-methoxyphenyl)- (CA INDEX NAME)



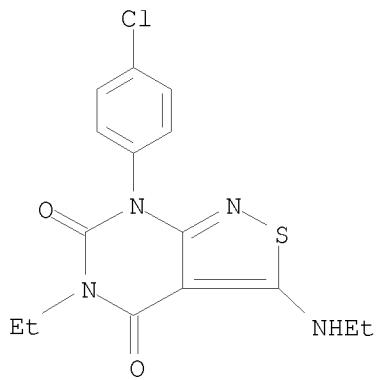
RN 60663-87-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylamino)-5-methyl-7-(phenylmethyl)- (CA INDEX NAME)



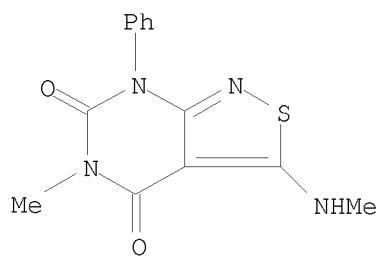
RN 60663-88-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-5-ethyl-3-(ethylamino)- (CA INDEX NAME)



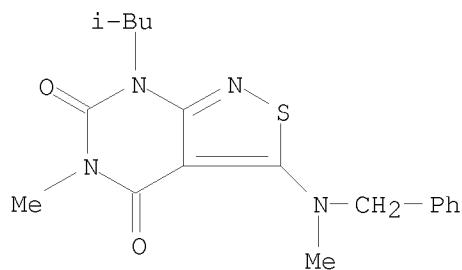
RN 60663-89-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-3-(methylamino)-7-phenyl- (CA INDEX NAME)



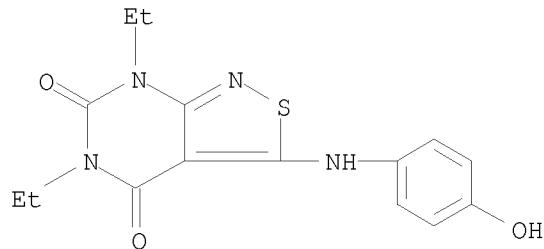
RN 60663-90-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5-methyl-3-[methyl(phenylmethyl)amino]-7-(2-methylpropyl)- (CA INDEX NAME)



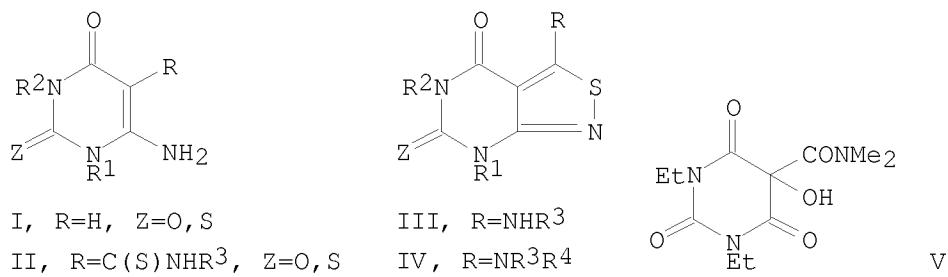
RN 60700-23-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-[(4-hydroxyphenyl)amino]-, monohydrobromide (9CI) (CA INDEX NAME)



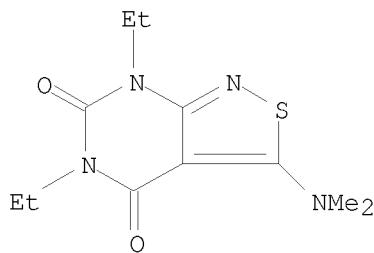
● HBr

GI

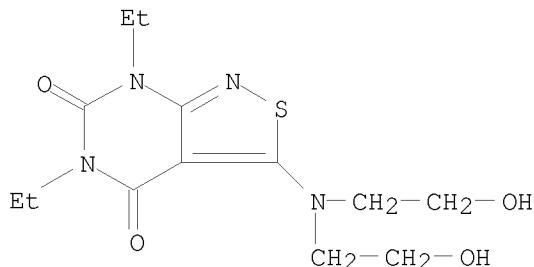


AB The 6-aminouracils I (R₁ = PhCH₂, Et, H, Me, p-ClC₆H₄, MeOCH₂CH₂, Ph, Me₂CHCH₂, p-MeOC₆H₄; R₂ = H, Et) reacted with SCNR₃ (R₃ = Et, Me, C₆H₄Cl-p, Ph, C₆H₄OH-p, CH₂Ph) to give the thiocarbamoyluracils II, which were oxidized with Br or H₂O₂ to give the isothiazolopyrimidinediones III, whose alkylation with R₄I (R₄ = Me, Et) gave the (disubstituted amino)isothiazolopyrimidines IV. Further oxidation of III or IV (R₁ = R₂ = Et, R₃ = R₄ = Me) with H₂O₂ gave the hydroxybarbituric acid V. Aminoisothiazolopyrimidines are potential cyclic nucleotide phosphodiesterase inhibitors.

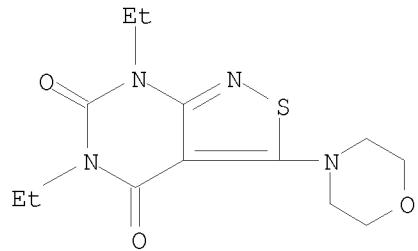
L6 ANSWER 18 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1976:560011 CAPLUS
 DOCUMENT NUMBER: 85:160011
 ORIGINAL REFERENCE NO.: 85:25613a,25616a
 TITLE: Fused pyrimidines. I. A one-step synthesis of 3-aminoisothiazolo[3,4-d]pyrimidines from 6-aminouracils and Vilsmeier reagents
 AUTHOR(S): Furukawa, Yoshiyasu; Miyashita, Osamu; Shima, Shunsuke
 CORPORATE SOURCE: Cent. Res. Div., Takeda Chem. Ind., Ltd., Osaka, Japan
 SOURCE: Chemical & Pharmaceutical Bulletin (1976), 24(5), 970-8
 CODEN: CPBTAL; ISSN: 0009-2363
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 85:160011
 IT 56540-86-4P 56540-88-6P 56540-89-7P
 56540-91-1P 56540-92-2P 56540-98-8P
 56540-99-9P 56541-00-5P 60663-76-5P
 60664-03-1P 60664-09-7P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 56540-86-4 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-diethyl- (CA INDEX NAME)



RN 56540-88-6 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[bis(2-hydroxyethyl)amino]-5,7-diethyl- (CA INDEX NAME)

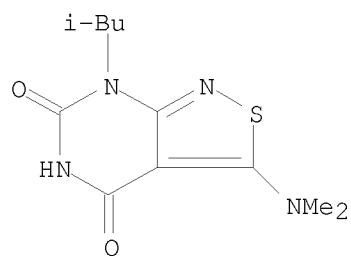


RN 56540-89-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-(4-morpholinyl)- (CA INDEX NAME)



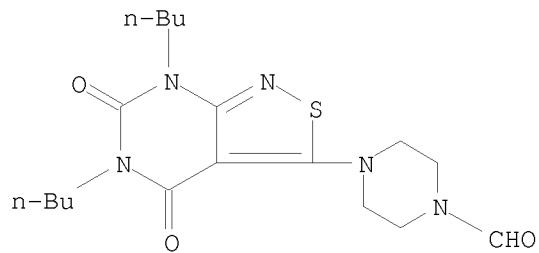
RN 56540-91-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-(2-methylpropyl)- (CA INDEX NAME)



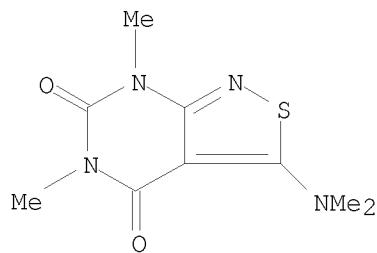
RN 56540-92-2 CAPLUS

CN 1-Piperazinecarboxaldehyde, 4-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)

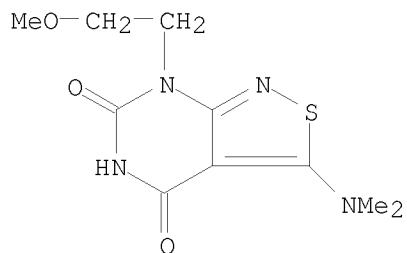


RN 56540-98-8 CAPLUS

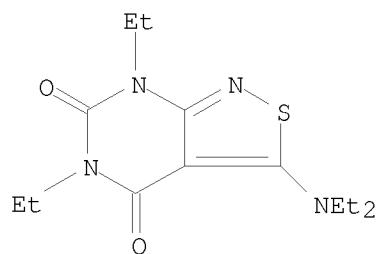
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-dimethyl- (CA INDEX NAME)



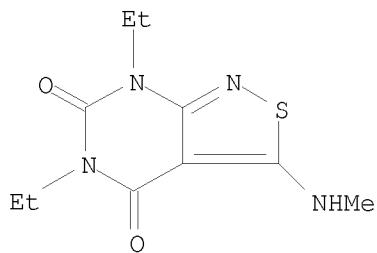
RN 56540-99-9 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-(2-methoxyethyl)- (CA INDEX NAME)



RN 56541-00-5 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(diethylamino)-5,7-diethyl- (CA INDEX NAME)

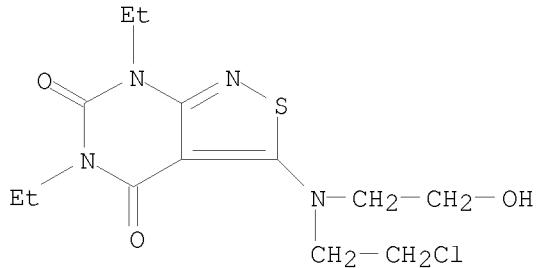


RN 60663-76-5 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-(methylamino)- (CA INDEX NAME)



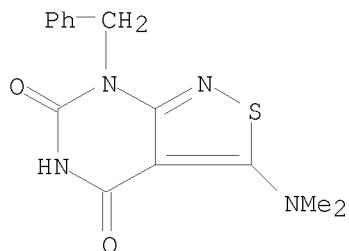
RN 60664-03-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(2-chloroethyl)(2-hydroxyethyl)amino]-5,7-diethyl- (CA INDEX NAME)

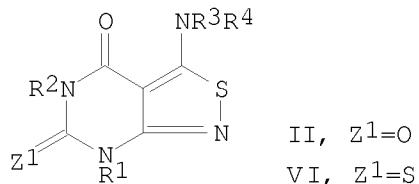
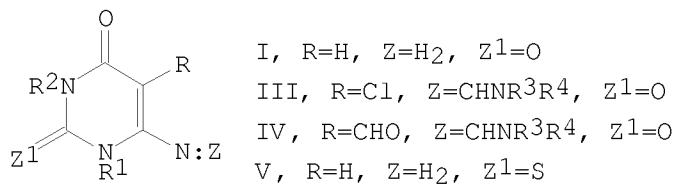


RN 60664-09-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-(phenylmethyl)- (CA INDEX NAME)



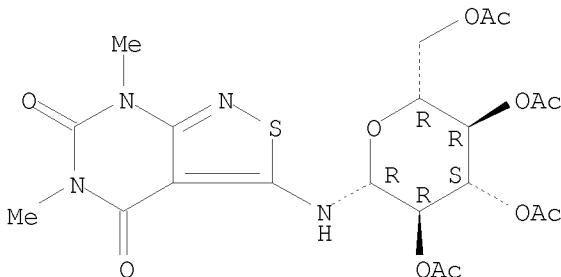
GI



AB Reaction of the aminouracil I (R₁ = R₂ = Et) with DMF-SOCl₂ gave the (dimethylamino)isothiazolopyrimidinedione II (R₃ = R₄ = Me) and the uracils III, IV, and V (R₃ = Me, R₄ = H) as minor products, whereas similar reactions of I (R₁ = R₂ = Et, Bu) or 6-amino-1-benzylcytocene with OHCNR₃R₄ (R₃, R₄ = CH₂CH₂OCHO, Et, or NR₃R₄ = morpholino, 4-formyl-1-piperazinyl)-SOCl₂ gave the corresponding II (R₁ = Et, Bu, CH₂Ph; R₂-R₄ = same as above). However, the S analog V (R₁ = R₂ = Et) reacted with DMF-SOCl₂ to give mainly II (R₁ = R₂ = Et, R₃ = R₄ = Me) and very little its S analog VI (R₁-R₄ same as above). Aminoisothiazolopyrimidines are potential cyclic nucleotide phosphodiesterase inhibitors.

L6 ANSWER 19 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1976:150881 CAPLUS
 DOCUMENT NUMBER: 84:150881
 ORIGINAL REFERENCE NO.: 84:24535a,24538a
 TITLE: Modified nucleoside syntheses
 AUTHOR(S): Ogura, Haruo; Takahashi, Hiroshi; Takeda, Kazuyoshi;
 Sakaguchi, Masakazu; Nimura, Noriyuki; Sakai, Hitomi
 CORPORATE SOURCE: Sch. Pharm. Sci., Kitasato Univ., Tokyo, Japan
 SOURCE: Hukusukan Kagaku Toronkai Koen Yoshishu, 8th (1975),
 154-8. Pharm. Inst., Tohoku Univ.: Sendai, Japan.
 CODEN: 32KOAD
 DOCUMENT TYPE: Conference
 LANGUAGE: Japanese
 IT 58911-57-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 58911-57-2 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl)amino]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



GI For diagram(s), see printed CA Issue.
 AB The isothiocyanates I and II were treated with $\text{o-C}_6\text{H}_4(\text{NH}_2)_2$ or 5,6-diamino-1,3-dimethyluracil to give the modified nucleosides III and IV ($\text{X} = \text{CH:CHCH:CH}$, NMeCONMeCO , resp.). IV reacted with MeI with elimination of MeSH and gave the corresponding imidazoles V. $\text{R}_2\text{COCH:C}(\text{NH}_2)\text{NMeR}_3$ ($\text{R}_2 = \text{EtO}$, $\text{R}_3 = \text{Me}$; $\text{R}_2\text{R}_3 = \text{NMeCONMe}$) reacted with I and II to give the corresponding pyrimidines (VI, VII, resp.).

L6 ANSWER 20 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1975:497362 CAPLUS
 DOCUMENT NUMBER: 83:97362
 ORIGINAL REFERENCE NO.: 83:15309a,15312a
 TITLE: 3-Disubstituted aminoisothiazolo[3,4-d]pyrimidines
 INVENTOR(S): Furukawa, Yoshiyasu; Miyashita, Osamu
 PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan
 SOURCE: Ger. Offen., 24 pp.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

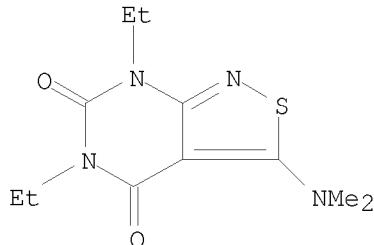
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2453212	A1	19750515	DE 1974-2453212	19741109
JP 50082092	A	19750703	JP 1973-127050	19731112
JP 58026356	B	19830602		
FR 2250532	A1	19750606	FR 1974-37112	19741108
FR 2250532	B1	19790504		
US 3959280	A	19760525	US 1974-522863	19741111
GB 1436297	A	19760519	GB 1974-48874	19741112
US 4052391	A	19771004	US 1976-663227	19760303
PRIORITY APPLN. INFO.:			JP 1973-127050	A 19731112
			US 1974-522863	A3 19741111

IT 56540-86-4P 56540-88-6P 56540-89-7P
 56540-90-0P 56540-91-1P 56540-92-2P
 56540-93-3P 56540-98-8P 56540-99-9P
 56541-00-5P 56541-01-6P 56541-02-7P
 56541-03-8P 56541-04-9P 56541-05-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

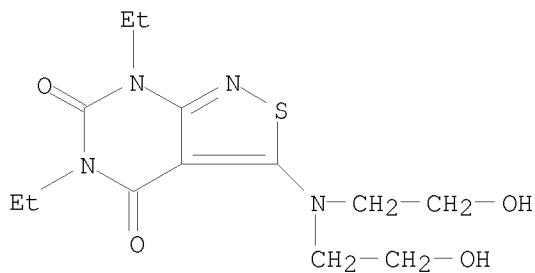
RN 56540-86-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-diethyl- (CA INDEX NAME)



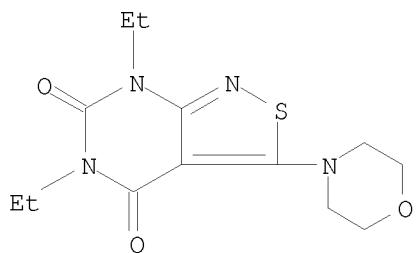
RN 56540-88-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[bis(2-hydroxyethyl)amino]-5,7-diethyl- (CA INDEX NAME)



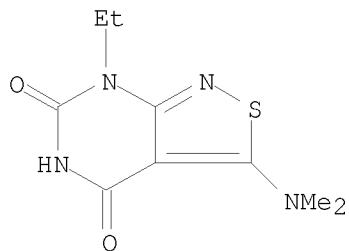
RN 56540-89-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-diethyl-3-(4-morpholiny)- (CA INDEX NAME)



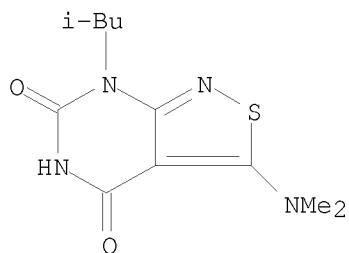
RN 56540-90-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-ethyl- (CA INDEX NAME)



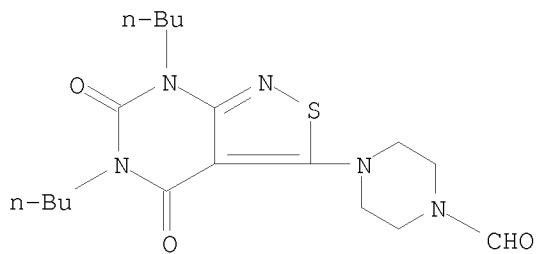
RN 56540-91-1 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-(2-methylpropyl)- (CA INDEX NAME)



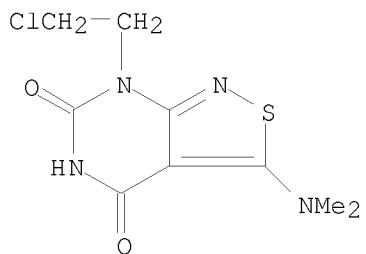
RN 56540-92-2 CAPLUS

CN 1-Piperazinecarboxaldehyde, 4-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



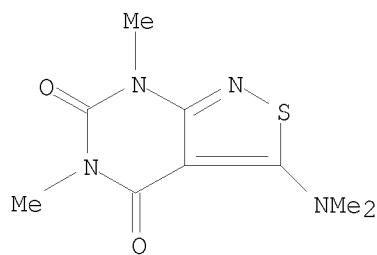
RN 56540-93-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-chloroethyl)-3-(dimethylamino)- (CA INDEX NAME)

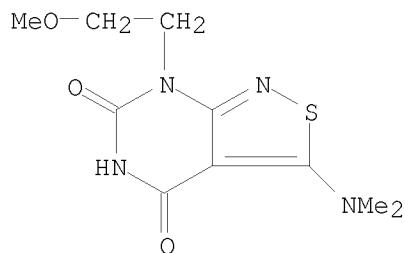


RN 56540-98-8 CAPLUS

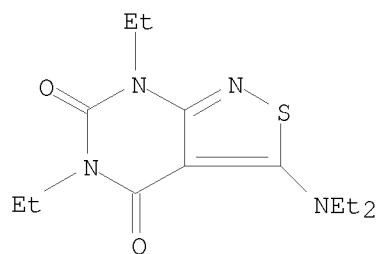
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5,7-dimethyl- (CA INDEX NAME)



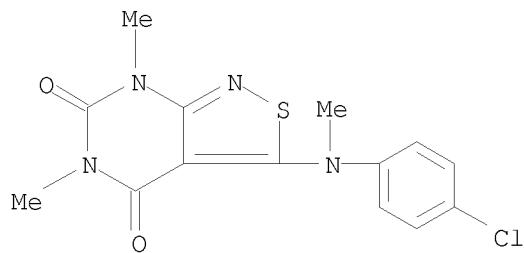
RN 56540-99-9 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-7-(2-methoxyethyl)- (CA INDEX NAME)



RN 56541-00-5 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(diethylamino)-5,7-diethyl- (CA INDEX NAME)

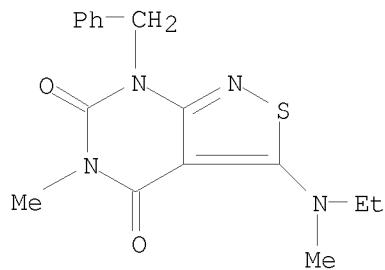


RN 56541-01-6 CAPLUS
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[(4-chlorophenyl)methylamino]-5,7-dimethyl- (CA INDEX NAME)



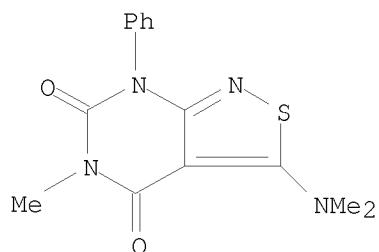
RN 56541-02-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(ethylmethylamino)-5-methyl-7-(phenylmethyl)- (CA INDEX NAME)



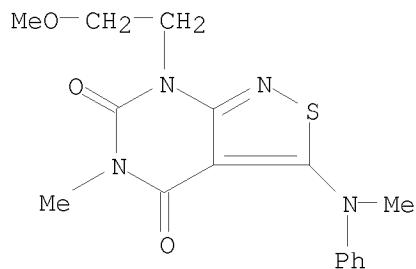
RN 56541-03-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-(dimethylamino)-5-methyl-7-phenyl- (CA INDEX NAME)



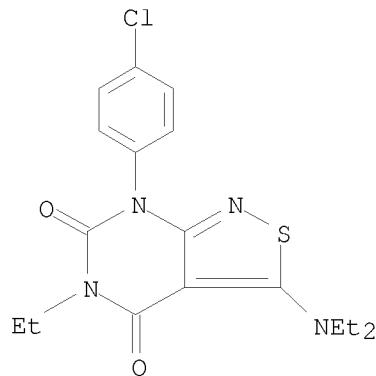
RN 56541-04-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(2-methoxyethyl)-5-methyl-3-(methylphenylamino)- (CA INDEX NAME)



RN 56541-05-0 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 7-(4-chlorophenyl)-3-(diethylamino)-5-ethyl- (CA INDEX NAME)

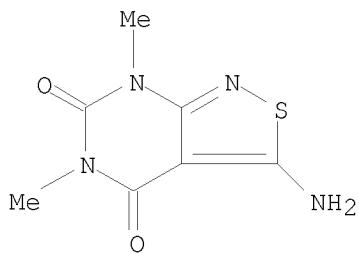


GI For diagram(s), see printed CA Issue.

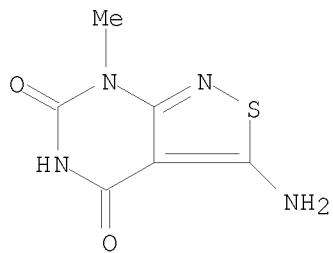
AB Isothiazolopyrimidines (I, R1 = Me, HOCH2CH2, Et, R2 = Me, HOCH2CH2, Et, p-ClC6H4, Ph, R3 = Et, Me2CHCH2, Bu, ClCH2CH2, Me, MeOCH2CH2, PhCH2, Ph, p-ClC6H4, R4 = Et, H, Bu, Me, R1R2N = morpholino, 4-formylpiperazinyl), useful as sedatives and inflammation inhibitors, were prepared by cyclization of an aminouracil (II) with R1R1NCHO. I were also useful as adenosine 3',5'-cyclicphosphate phosphodiesterase inhibitors.

L6 ANSWER 21 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1975:412166 CAPLUS
 DOCUMENT NUMBER: 83:12166
 ORIGINAL REFERENCE NO.: 83:2035a,2038a
 TITLE: Isothiazolopyrimidindione dyes
 INVENTOR(S): Eilingsfeld, Heinz; Hanse, Guenter
 PATENT ASSIGNEE(S): BASF A.-G.
 SOURCE: Ger. Offen., 30 pp.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2336978	A1	19750206	DE 1973-2336978	19730720
PRIORITY APPLN. INFO.:			DE 1973-2336978	A 19730720
IT 52903-30-7				
RL: USES (Uses)				
(coupling of diazotized, with (hydroxyethyl)ethylaniline)				
RN 52903-30-7 CAPLUS				
CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dimethyl- (CA INDEX NAME)				

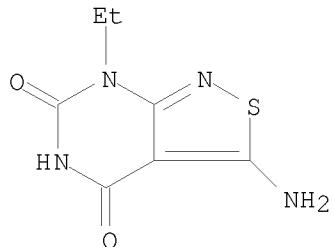


IT 52903-28-3 52903-29-4 52903-32-9
 RL: USES (Uses)
 (coupling of diazotized, with [(cyanoethyl)ethylamino]benzene)
 RN 52903-28-3 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-methyl- (CA INDEX NAME)



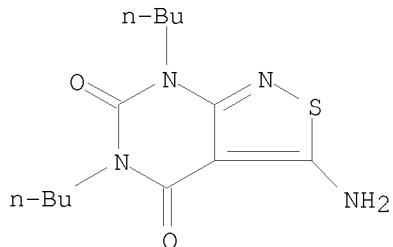
RN 52903-29-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-ethyl- (CA INDEX NAME)



RN 52903-32-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dibutyl- (CA INDEX NAME)



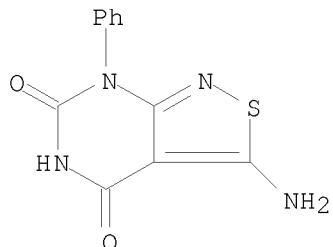
IT 52956-48-6

RL: USES (Uses)

(coupling with diazotized, with (diethylamino)acetanalide)

RN 52956-48-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-phenyl- (CA INDEX NAME)



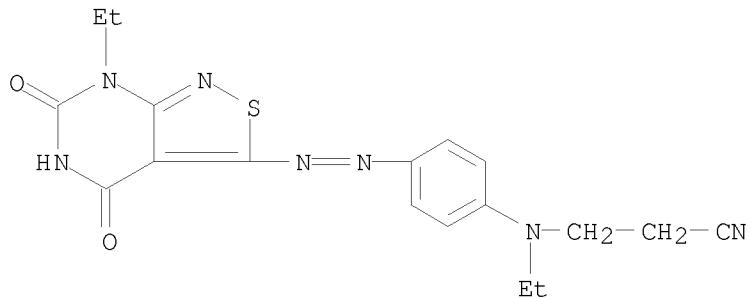
IT 55345-46-5P

RL: IMF (Industrial manufacture); PREP (Preparation)

(preparation and acetate and polyester fiber dyeing by)

RN 55345-46-5 CAPLUS

CN Propanenitrile, 3-[ethyl[4-[(7-ethyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azolphenyl]amino]- (9CI) (CA INDEX NAME)



IT 55345-45-4P 55345-47-6P 55345-49-8P

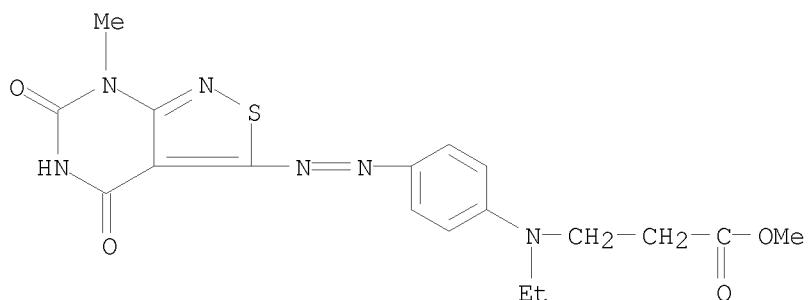
55345-51-2P

RL: IMF (Industrial manufacture); PREP (Preparation)

(preparation and acetate, polyamide and polyester fiber dyeing by)

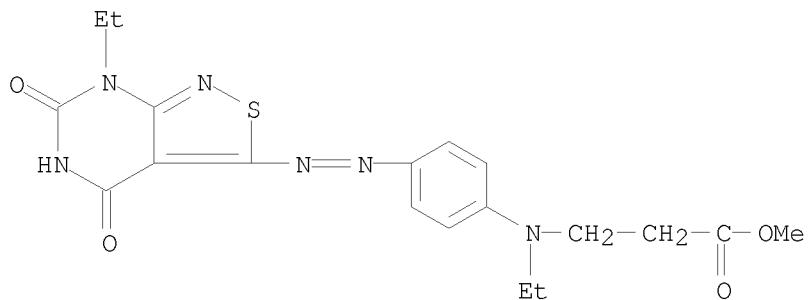
RN 55345-45-4 CAPPLUS

CN β -Alanine, N-ethyl-N-[4-[(4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azo]phenyl]-, methyl ester (9CI)
(CA INDEX NAME)



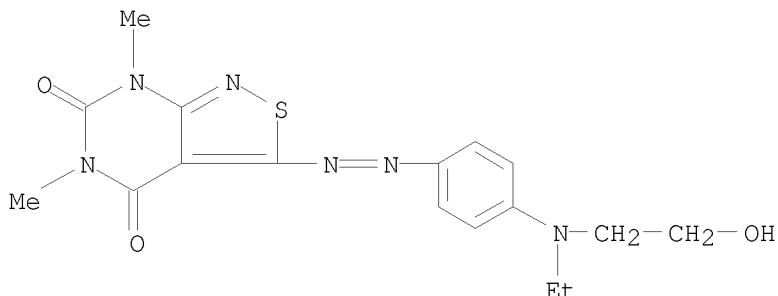
RN 55345-47-6 CAPPLUS

CN β -Alanine, N-ethyl-N-[4-[(7-ethyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azo]phenyl]-, methyl ester (9CI)
(CA INDEX NAME)

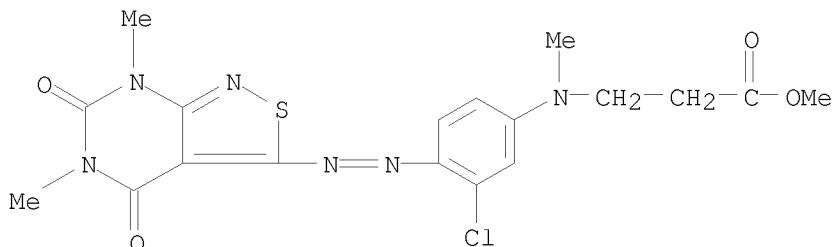


RN 55345-49-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-[[4-[ethyl(2-hydroxyethyl)amino]phenyl]azo]-5,7-dimethyl- (9CI) (CA INDEX NAME)



RN 55345-51-2 CAPLUS

CN β -Alanine, N-[3-chloro-4-[(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azo]phenyl]-N-methyl-, methyl ester (9CI) (CA INDEX NAME)

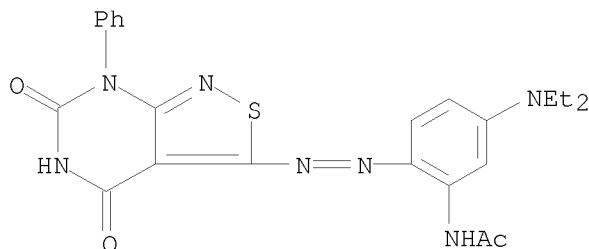
IT 55345-48-7P 55345-52-3P 55345-53-4P

55345-54-5P

RL: IMF (Industrial manufacture); PREP (Preparation)
(preparation and polyester fiber dyeing by)

RN 55345-48-7 CAPLUS

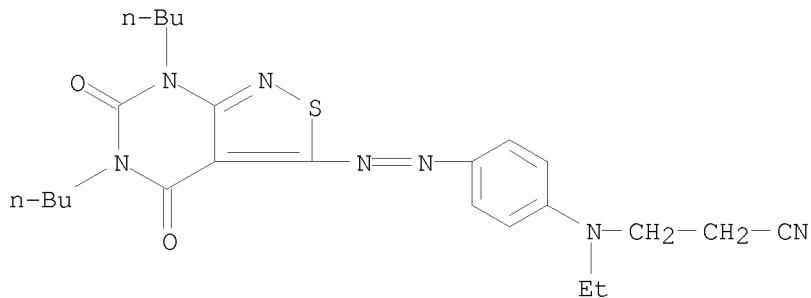
CN Acetamide, N-[5-(diethylamino)-2-[(4,5,6,7-tetrahydro-4,6-dioxo-7-phenylisothiazolo[3,4-d]pyrimidin-3-yl)azo]phenyl]- (9CI) (CA INDEX NAME)



RN 55345-52-3 CAPLUS

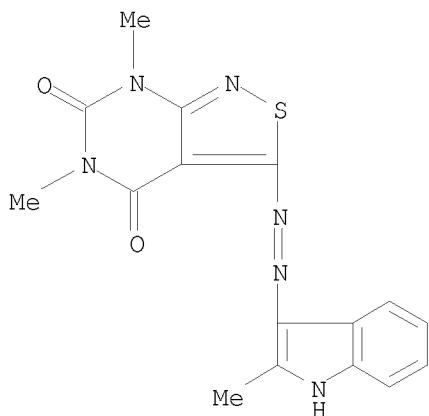
CN Propanenitrile, 3-[[4-[(5,7-dibutyl-4,5,6,7-tetrahydro-4,

dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azolphenyl]ethylamino]- (9CI) (CA INDEX NAME)



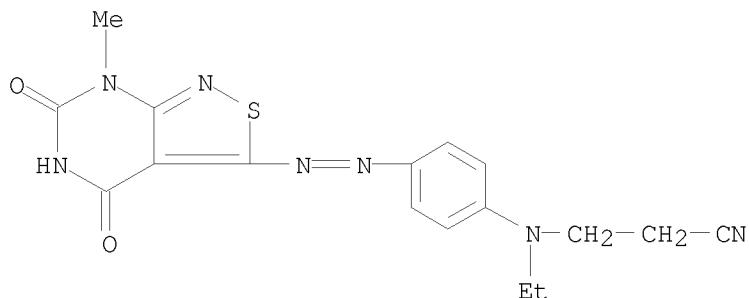
RN 55345-53-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 5,7-dimethyl-3-[(2-methyl-1H-indol-3-yl)azo]- (9CI) (CA INDEX NAME)



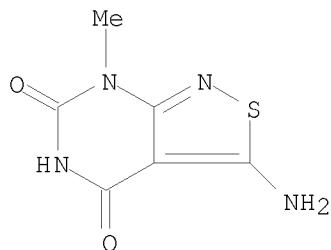
RN 55345-54-5 CAPLUS

CN Propanenitrile, 3-[ethyl[4-[(4,5,6,7-tetrahydro-7-methyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)azolphenyl]amino]- (9CI) (CA INDEX NAME)

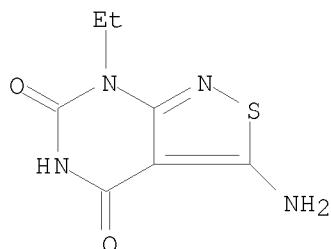


GI For diagram(s), see printed CA Issue.
AB Isothiazolopyrimidine azo dyes (I, R = H, Pr, Bu; R1 = Et, Ph, Pr, Bu; R2 = N,N-disubstituted p-aniline derivs., 2-methyl-3-indolyl) were prepared and were used to dye polyester, polyamide, and acetate fibers orange to violet shades. Thus, 3-amino-7-methylisothiazolo[3,4-d]pyrimidine-4,6-dione [52903-28-3] was diazotized and coupled with PhNETCH₂CH₂CN [148-87-8] to give azo dye (II) [55345-54-5], bordeaux on polyester fibers. The other I were similarly prepared

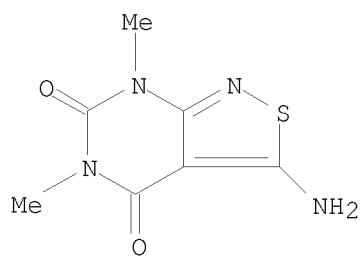
L6 ANSWER 22 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1975:170825 CAPLUS
 DOCUMENT NUMBER: 82:170825
 ORIGINAL REFERENCE NO.: 82:27301a,27304a
 TITLE: Isothiazolo[3,4-d]- and pyrimido[4,5-d]-pyrimidines
 AUTHOR(S): Niess, Rolf; Eilingsfeld, Heinz
 CORPORATE SOURCE: Farbenlab., BASF A.-G., Ludwigshafen, Fed. Rep. Ger.
 SOURCE: Justus Liebigs Annalen der Chemie (1975), Volume Date
 1974, (12), 2019-29
 CODEN: JLACBF; ISSN: 0075-4617
 DOCUMENT TYPE: Journal
 LANGUAGE: German
 OTHER SOURCE(S): CASREACT 82:170825
 IT 52903-28-3P 52903-29-4P 52903-30-7P
 52903-31-8P 52903-32-9P 52903-33-0P
 52903-35-2P 52903-36-3P 52903-37-4P
 52903-38-5P 52903-40-9P 52903-41-0P
 52903-42-1P 52956-48-6P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 52903-28-3 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-methyl- (CA INDEX NAME)



RN 52903-29-4 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-ethyl- (CA INDEX NAME)

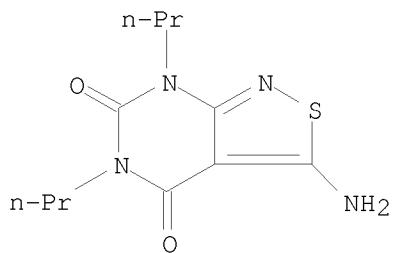


RN 52903-30-7 CAPLUS
 CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dimethyl- (CA INDEX NAME)



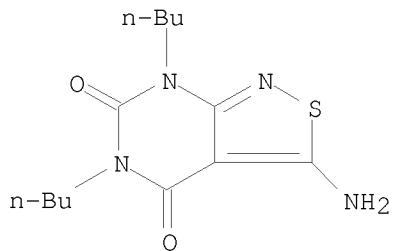
RN 52903-31-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dipropyl- (CA INDEX NAME)



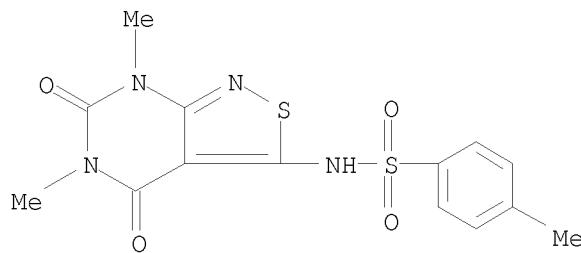
RN 52903-32-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dibutyl- (CA INDEX NAME)

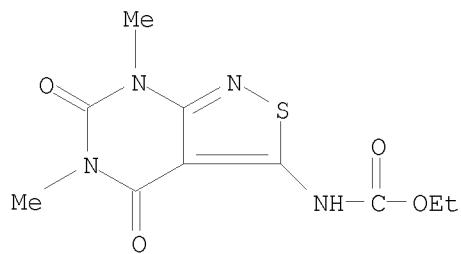


RN 52903-33-0 CAPLUS

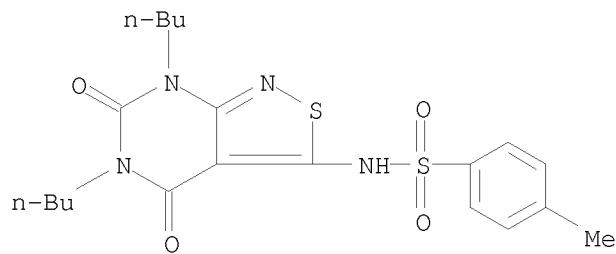
CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



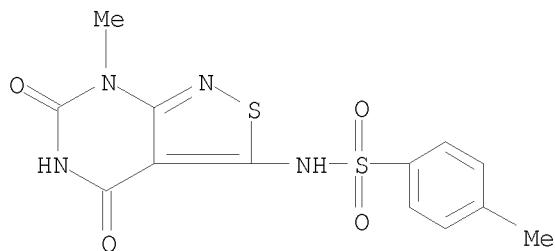
RN 52903-35-2 CAPLUS
CN Carbamic acid, (4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)-, ethyl ester (9CI) (CA INDEX NAME)



RN 52903-36-3 CAPLUS
CN Benzenesulfonamide, N-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)-4-methyl- (CA INDEX NAME)

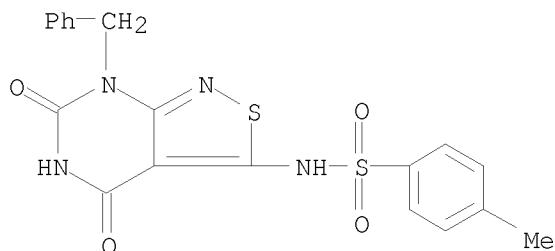


RN 52903-37-4 CAPLUS
CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-7-methyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



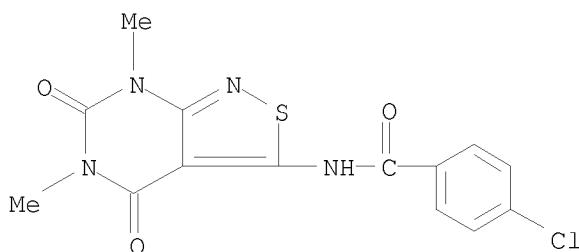
RN 52903-38-5 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-[4,5,6,7-tetrahydro-4,6-dioxo-7-(phenylmethyl)isothiazolo[3,4-d]pyrimidin-3-yl]- (CA INDEX NAME)



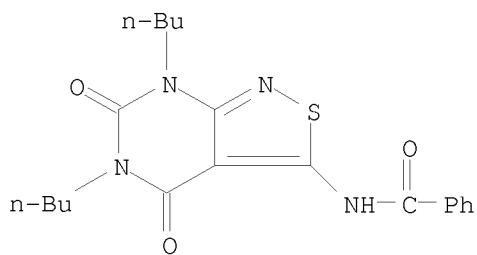
RN 52903-40-9 CAPLUS

CN Benzamide, 4-chloro-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



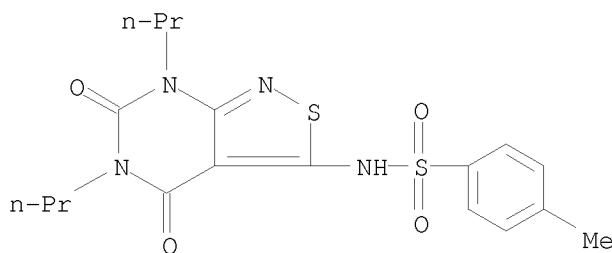
RN 52903-41-0 CAPLUS

CN Benzamide, N-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



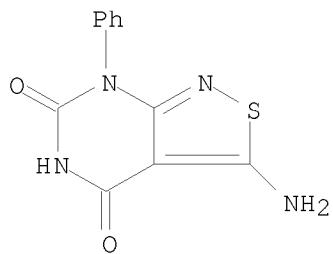
RN 52903-42-1 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-4,6-dioxo-5,7-dipropylisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



RN 52956-48-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-phenyl- (CA INDEX NAME)

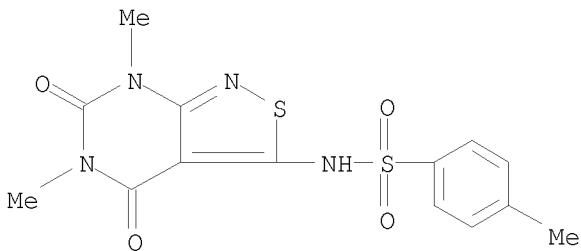


GI For diagram(s), see printed CA Issue.

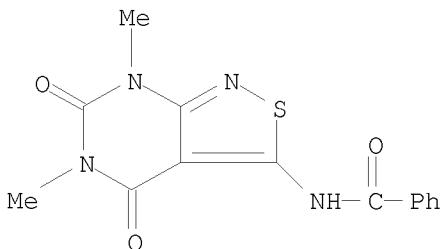
AB The isothiazolopyrimidines I (R = e.g. SMe, SCH₂Ph, NMe₂, morpholino, or Ph) and II (R₁ = e.g. H or Pr, R₂ = e.g. Me or Bu, R₃ = e.g. H, SO₂C₆H₄Me-4, or CO₂Et) were prepared by oxidative cyclization of the corresponding carboxamides III and IV (R₄ = CSNHR₃), resp. IV were prepared by reaction of IV (R₄ = H) (V) with R₃NCS. Reaction of V with R₅CONCS (R₅ = e.g. 4-O₂NC₆H₄) and cyclization gave the derivs. VI.

L6 ANSWER 23 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1974:413488 CAPLUS
 DOCUMENT NUMBER: 81:13488
 ORIGINAL REFERENCE NO.: 81:2171a,2174a
 TITLE: 3-Aminoisothiazolo[3,4-d]pyrimidines
 INVENTOR(S): Niess, Rolf; Eilingsfeld, Heinz
 PATENT ASSIGNEE(S): BASF A.-G.
 SOURCE: Ger. Offen., 19 pp.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2248231	A1	19740411	DE 1972-2248231	19721002
CH 578008	A5	19760730	CH 1973-13917	19730928
FR 2201097	A1	19740426	FR 1973-35018	19731001
GB 1445697	A	19760811	GB 1973-45725	19731001
PRIORITY APPLN. INFO.:			DE 1972-2248231	A 19721002
IT 52903-33-0	52903-34-1	52903-35-2		
52903-36-3				
RL: RCT (Reactant); RACT (Reactant or reagent) (hydrolysis of)				
RN 52903-33-0	CAPLUS			
CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)				

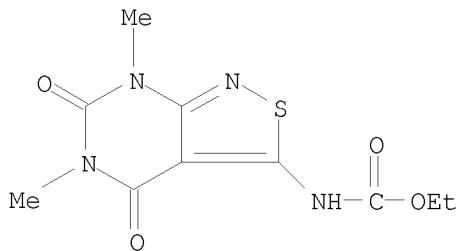


RN 52903-34-1 CAPLUS
 CN Benzenesulfonamide, N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



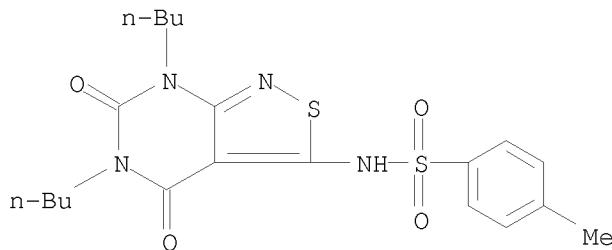
RN 52903-35-2 CAPLUS

CN Carbamic acid, (4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)-, ethyl ester (9CI) (CA INDEX NAME)



RN 52903-36-3 CAPLUS

CN Benzenesulfonamide, N-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)-4-methyl- (CA INDEX NAME)



IT 52903-28-3P 52903-29-4P 52903-30-7P

52903-31-8P 52903-32-9P 52903-36-3P

52903-37-4P 52903-38-5P 52903-39-6P

52903-40-9P 52903-41-0P 52903-42-1P

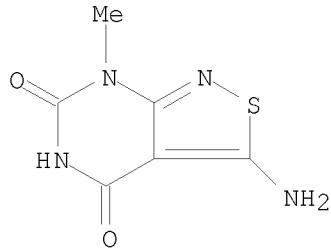
52903-43-2P 52903-44-3P 52956-48-6P

52956-49-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

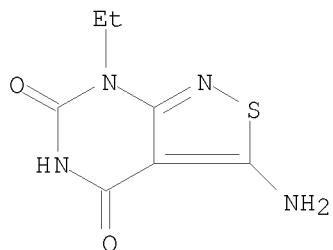
RN 52903-28-3 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-methyl- (CA INDEX NAME)



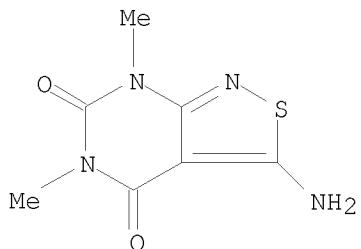
RN 52903-29-4 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-ethyl- (CA INDEX NAME)



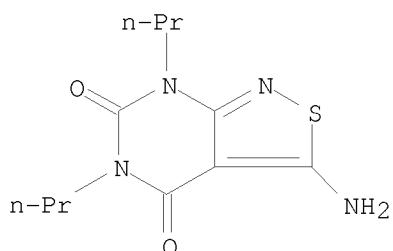
RN 52903-30-7 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dimethyl- (CA INDEX NAME)



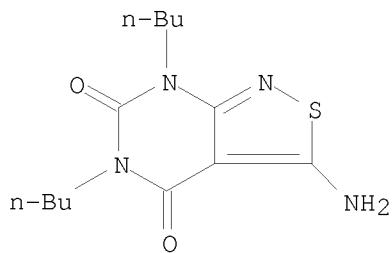
RN 52903-31-8 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dipropyl- (CA INDEX NAME)



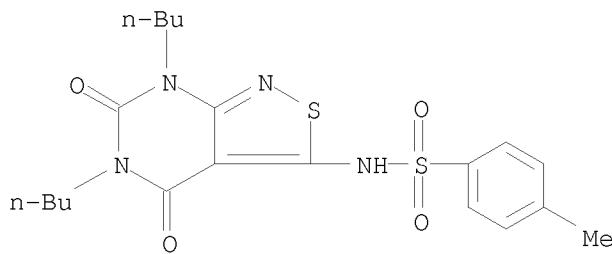
RN 52903-32-9 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-5,7-dibutyl- (CA INDEX NAME)



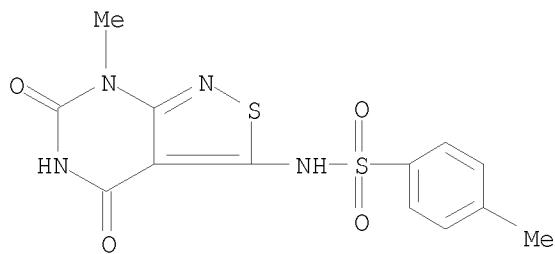
RN 52903-36-3 CAPLUS

CN Benzenesulfonamide, N-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)-4-methyl- (CA INDEX NAME)



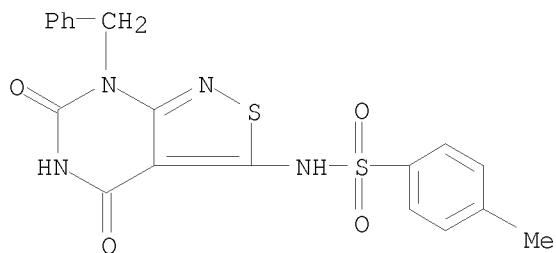
RN 52903-37-4 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-7-methyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



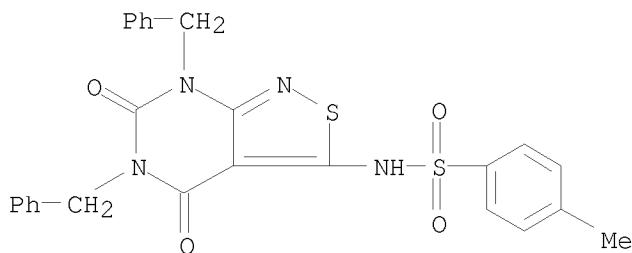
RN 52903-38-5 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-[4,5,6,7-tetrahydro-4,6-dioxo-7-(phenylmethyl)isothiazolo[3,4-d]pyrimidin-3-yl]- (CA INDEX NAME)



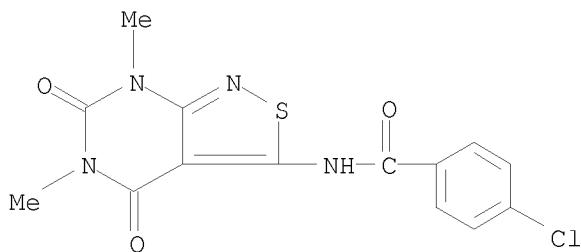
RN 52903-39-6 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-[4,5,6,7-tetrahydro-4,6-dioxo-5,7-bis(phenylmethyl)isothiazolo[3,4-d]pyrimidin-3-yl]- (CA INDEX NAME)



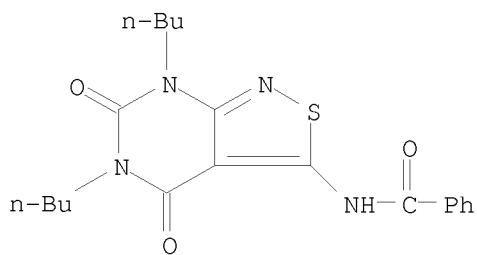
RN 52903-40-9 CAPLUS

CN Benzamide, 4-chloro-N-(4,5,6,7-tetrahydro-5,7-dimethyl-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



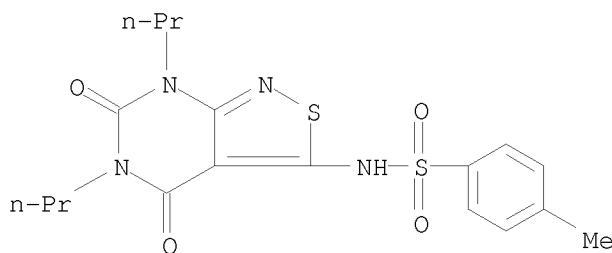
RN 52903-41-0 CAPLUS

CN Benzamide, N-(5,7-dibutyl-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



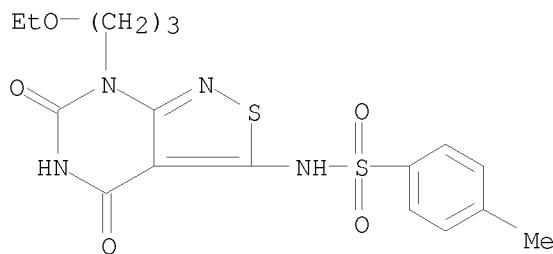
RN 52903-42-1 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-(4,5,6,7-tetrahydro-4,6-dioxo-5,7-dipropylisothiazolo[3,4-d]pyrimidin-3-yl)- (CA INDEX NAME)



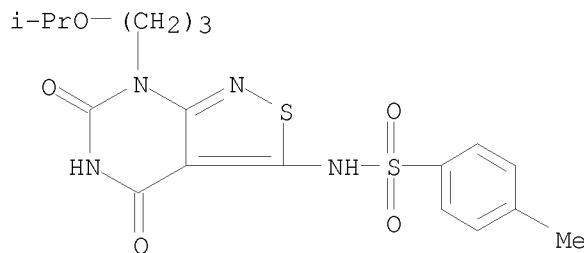
RN 52903-43-2 CAPLUS

CN Benzenesulfonamide, N-[7-(3-ethoxypropyl)-4,5,6,7-tetrahydro-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl]-4-methyl- (CA INDEX NAME)



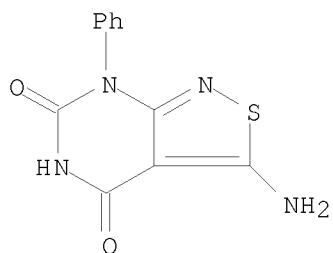
RN 52903-44-3 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-[4,5,6,7-tetrahydro-7-[3-(1-methylethoxy)propyl]-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl]- (CA INDEX NAME)



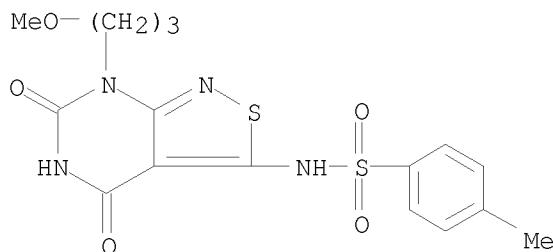
RN 52956-48-6 CAPLUS

CN Isothiazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione, 3-amino-7-phenyl- (CA INDEX NAME)



RN 52956-49-7 CAPLUS

CN Benzenesulfonamide, 4-methyl-N-[4,5,6,7-tetrahydro-7-(3-methoxypropyl)-4,6-dioxoisothiazolo[3,4-d]pyrimidin-3-yl]- (CA INDEX NAME)



GI For diagram(s), see printed CA Issue.

AB Isothiazolopyrimidines I (R = H, Me, Bu, Pr, CH₂Ph; R₁ = Me, Et, Pr, Bu, Ph, CH₂Ph, (CH₂)₃OMe, (CH₂)₃OEt, (CH₂)₃-OCHMe₂; R₂ = H, SO₂C₆H₄Me-p; R = R₁ = Me, R₂ = CO₂-Et, COC₆H₄Cl-p, Bz, SO₂C₆H₄Me-p) were prepared by cyclizing I (R₂ ≠ H), which may be accompanied by hydrolysis of R₂. Thus, 88% I (R = R₂ = H, R₁ = Me) was obtained by cyclizing II (R = H, R₁ = Me, R₂ = SO₂C₆H₄Me-p) with H₂SO₄. Some I, especially I (R = R₁ = Me, R₂ = H, CO₂Et; R = R₁ = Pr, R₂ = H) had diuretic and antiinflammatory activities.

10/141,986

=>

---Logging off of STN---

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Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	143.75	322.74
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-18.40	-18.40

STN INTERNATIONAL LOGOFF AT 15:46:39 ON 03 JUL 2008